

Search Report

STIC Database Tracking Notice

To: KUO-LIANG PENG Location: REM-10A71

Art Unit: 1712

Wednesday, July 11, 2007

Case Serial Number: 10/520763

From: KATHLEEN FULLER

Location: EIC1700

REM-4B28 / REM-4B19 Phone: (571)272-2505

kathleen.fuller@uspto.gov

Search Notes		
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ANSWER 1 OF 1 HCAPLUS COPYRIGHT 2007 ACS on STN
L1
     2004:60574 HCAPLUS
AN
     140:94920
DN
     Entered STN: 26 Jan 2004
ED
     Hybrid polymers for functional tuning of microfluidic device surfaces
ΤI
     Augustine, Brian H.; Landers, James P.; Ferrance, Jerome P.; Polefrone,
IN
     Joy; Hugues, W. Christopher
     University of Virginia Patent Foundation, USA; James Madison University
PA
                                                           application - not
indeped with
     PCT Int. Appl., 43 pp.
SO
     CODEN: PIXXD2
DT
     Patent
LΑ
     English
     ICM CO8G
IC
     37-3 (Plastics Manufacture and Processing)
CC
FAN.CNT 1
                          KIND
                                             APPLICATION NO.
                                                                       DATE
     PATENT NO.
                                  DATE
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                          ----
                                 -----
                                              -----
                                              WO 2003-US22162
                                                                       20030715
     WO 2004007582
                           A2
                                  20040122
PΙ
     WO 2004007582
                          A3
                                  20040325
             AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
              CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
              GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
              LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM,
              PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT,
              TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
              KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
                          A1 20040202
                                           AU 2003-251935
     AU 2003251935
                                                                20030715
                           A1
                                  20060316
                                              US 2005-520763
                                                                      20050110 <--
     US 2006057402
                           Ρ
PRAI US 2002-396153P
                                  20020715
     US 2002-399633P
                           Ρ
                                  20020730
     WO 2003-US22162
                                  20030715
CLASS
                  CLASS PATENT FAMILY CLASSIFICATION CODES
 PATENT NO.
                         C08G
 WO 2004007582
                  ICM
                  IPCI
                         C08G [ICM, 7]
                         B01L0003-00 [N,C*]; B01L0003-00 [N,A]; G01N0027-447
                  IPCR
                         [I,C*]; G01N0027-447 [I,A]
                  ECLA
                         G01N027/447B6
 AU 2003251935
                  IPCI
                         B32B0017-06 [ICM,7]; B32B0025-20 [ICS,7]; B32B0025-00
                         [ICS,7,C*]; C09J0143-04 [ICS,7]; C09J0143-00 [ICS,7,C*]
                  IPCR
                         B01L0003-00 [N,C*]; B01L0003-00 [N,A]; G01N0027-447
                         [I,C*]; G01N0027-447 [I,A]
                         B32B0009-04 [I,A]
 US 2006057402
                  IPCI
                  IPCR
                         B32B0009-04 [I,A]; B32B0009-04 [I,C]
                  NCL
                         428/447.000
     A microfluidic device comprises a body structure provided with a
AB
     microchannel and an inlet port and an outlet port, wherein the inlet port
     and outlet port are formed on an exterior surface of the body structure
     and are in fluid communication with the microchannel, wherein the
     microchannel has an interior surface that is coated with a polymer
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comprising Me methacrylate repeating units and acrylate derivs. of

polyhedral oligomeric silsesquioxanes. In addition the polymer can be used to coat microchannels to enhance the phys. properties of the microdevice.

- ST microfluidic device POSS MMA polymer coating; silsesquioxane acrylate polymer coating microfluidic device
- IT Apparatus

(microfluidic devices; hybrid polymers for functional tuning of microfluidic device surfaces)

- IT Glass, uses
 - Polycarbonates, uses
 - Polyesters, uses
 - RL: DEV (Device component use); USES (Uses) (substrate; hybrid polymers for functional tuning of microfluidic device surfaces)
- IT Plastics, uses
 - RL: DEV (Device component use); USES (Uses) (thermoplastics, substrate; hybrid polymers for functional tuning of microfluidic device surfaces)
- IT 80-62-6D, MMA, polymers with polyhedral oligomeric silsesquioxanes
 RL: POF (Polymer in formulation); TEM (Technical or engineered material
 use); USES (Uses)
 - (hybrid polymers for functional tuning of microfluidic device surfaces)
- IT 7631-86-9, Silica, uses 9011-14-7, PMMA 25038-59-9, Polyethylene terephthalate, uses
 - RL: DEV (Device component use); USES (Uses)
 (substrate; hybrid polymers for functional tuning of microfluidic
 device surfaces)

=>

S E1-E4

1 25038-59-9/BI (25038-59-9/RN) 1 7631-86-9/BI (7631-86-9/RN) 1 80-62-6/BI (80-62-6/RN) 1 9011-14-7/BI

(9011-14-7/RN)

L2 4 (25038-59-9/BI OR 7631-86-9/BI OR 80-62-6/BI OR 9011-14-7/BI)

=> D SCAN

L2 4 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN IN Silica ADDITIONAL NAMES NOT AVAILABLE IN THIS FORMAT MF O2 Si

o = si = o

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):3

L2 '4 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN 2-Propenoic acid, 2-methyl-, methyl ester, homopolymer
ADDITIONAL NAMES NOT AVAILABLE IN THIS FORMAT
MF (C5 H8 O2)x
CI PMS, COM

CM 1

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 4 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Poly(oxy-1,2-ethanediyloxycarbonyl-1,4-phenylenecarbonyl)
ADDITIONAL NAMES NOT AVAILABLE IN THIS FORMAT
MF (C10 H8 O4)n
CI PMS, COM

RELATED POLYMERS AVAILABLE WITH POLYLINK

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 4 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN

IN 2-Propenoic acid, 2-methyl-, methyl ester

MF C5 H8 O2

CI COM

$$\begin{array}{c|c} ^{H_2C} & \text{O} \\ \parallel & \parallel \\ \text{Me-} & \text{C-} & \text{C-} & \text{OMe} \end{array}$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

ALL ANSWERS HAVE BEEN SCANNED

=>

Structure Search

=> file reg
FILE 'REGISTRY' ENTERED AT 11:51:56 ON 11 JUL 2007
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
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Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 10 JUL 2007 HIGHEST RN 942116-98-5 DICTIONARY FILE UPDATES: 10 JUL 2007 HIGHEST RN 942116-98-5

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH December 2, 2006

Please note that search-term pricing does apply when conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

http://www.cas.org/support/stngen/stndoc/properties.html

=> fil hcapl
FILE 'HCAPLUS' ENTERED AT 11:52:03 ON 11 JUL 2007
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
PLEASE SEE "HELP USAGETERMS" FOR DETAILS.
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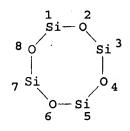
FILE COVERS 1907 - 11 Jul 2007 VOL 147 ISS 3 FILE LAST UPDATED: 10 Jul 2007 (20070710/ED)

New CAS Information Use Policies, enter HELP USAGETERMS for details.

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> d que 136

L3 STR



9,434 structures from query

NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS

STEREO ATTRIBUTES: NONE

L5 9434 SEA FILE=REGISTRY SSS FUL L3

L8 STR

5 0 |||| C~~ C~~ C~~ 0 1 2 3 4 subset search with

NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

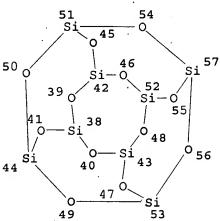
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NUMBER OF NODES IS 5

STEREO ATTRIBUTES: NONE

L11

STR



and

NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

361 Shudwes

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GRAPH ATTRIBUTES:
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RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 20

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STEREO ATTRIBUTES: NONE
L19
            361 SEA FILE=REGISTRY SUB=L5 SSS FUL L8 AND L11
            195 SEA FILE=HCAPLUS ABB=ON
L20
                                         L19
L21.
              1 SEA FILE=HCAPLUS ABB=ON L20 AND (MICRO(W)FLUID? OR MICROFLUID?
L22
            125 SEA FILE=HCAPLUS ABB=ON L20(L)PREP/RL
L23
              1 SEA FILE=HCAPLUS ABB=ON
                                         L22 AND TUNING
L24
              1 SEA FILE=HCAPLUS ABB=ON
                                         L19 AND TUNING
L25
              2 SEA FILE=HCAPLUS ABB=ON L21 OR L23 OR L24
L26
             49 SEA FILE=REGISTRY ABB=ON 2 7080.1.1/RID
L28
             41 SEA FILE=HCAPLUS ABB=ON L26
              2 SEA FILE=HCAPLUS ABB=ON L28 AND ?ACRYL?
L29
L30
             32 SEA FILE=HCAPLUS ABB=ON L28(L)PREP/RL
L31
              4 SEA FILE=HCAPLUS ABB=ON L25 OR L29
L33
              4 SEA FILE=HCAPLUS ABB=ON L30 AND HYBRID?
L34
              1 SEA FILE=HCAPLUS ABB=ON L30 AND TUN?
L35
              7 SEA FILE=HCAPLUS ABB=ON L20 AND TUN?
L36
             15 SEA FILE=HCAPLUS ABB=ON L31 OR L33 OR L34 OR L35
```

=> d l36 bib abs ind/hitstr 1-15

L36 ANSWER 1 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2007:372006 HCAPLUS

DN 146:501822

TI Filled polysilsesquioxanes: a new approach to chemical sensing

AU Castaldo, Anna; Massera, Ettore; Quercia, Luigi; Di Francia, Girolamo

CS ENEA Research Center of Portici UTS Mat-Nano, Granatello Portici, Italy

SO Macromolecular Symposia (2007), 247 (Times of Polymers and Composites),

CODEN: MSYMEC; ISSN: 1022-1360

PB Wiley-VCH Verlag GmbH & Co. KGaA

DT Journal

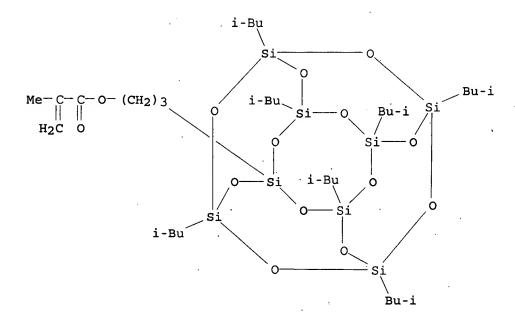
LA English

AB In an attempt to investigate new polymeric materials as constituents of an e-nose we focus our attention on a new emerging class of versatile three-dimensional polyhedral silicon polymers, called polysilsesquioxanes, containing nanosized inorg. cages. Such hybrid amphiphilic materials offer exceptional opportunities to create composites with singular properties. In particular we found that the polyhedral organosilsesquioxane (POSS) cages covalently attached to the polymer backbone as side-chain act as an "internal" filler with a nanometric homogeneous dispersion. We show how it is possible to fabricate sensing devices based on selected POSS matrix and, by a suitable choice of other "external" home-made fillers, (e.g. graphite, copper, silicon, zinc and their alloys) obtained by mech. milling the response of the resulting composites towards different classes of compds. can be tuned. In particular we fabricated a new high sensitive relative humidity device, exhibiting a fivefold response change for relative humidity changing in the range 0% to 100%. This behavior can not be modelled on the basis of the matrix swelling operating mechanism. Rather, the combined effect of the matrix and the filler has to be invoked in order to explain the formation of nanopores inside the material that are responsible of the porous behavior of our sensors.

CC 37-6 (Plastics Manufacture and Processing)

ST filled polysilsesquioxanes approach chem sensing graphite copper silicon zinc

```
IT
     Nanoparticles
        (filler; nanoparticles filled polysilsesquioxanes for to chemical sensing)
IT
     Fillers
     Polymer morphology
     Sensors
        (nanoparticles filled polysilsesquioxanes for to chemical sensing)
IT
     Silsesquioxanes
     RL: POF (Polymer in formulation); PRP (Properties); TEM (Technical or
     engineered material use); USES (Uses)
        (nanoparticles filled polysilsesquioxanes for to chemical sensing)
     7440-66-6, Zinc, uses 7782-42-5, Graphite, uses
IT
                                                        59873-66-4.
     Copper-Graphite 69704-06-9, Zinc Graphite
     RL: MOA (Modifier or additive use); USES (Uses)
        (conductive filler; nanoparticles filled polysilsesquioxanes for to
        chemical sensing)
IT
     37382-33-5, Graphite silicide
     RL: MOA (Modifier or additive use); USES (Uses)
        (filler; nanoparticles filled polysilsesquioxanes for to chemical sensing)
IT
     7440-21-3, Silicon, uses
                                7440-50-8, Copper, uses
     RL: MOA (Modifier or additive use); USES (Uses)
        (nanoparticles filled polysilsesquioxanes for to chemical sensing)
IT
     936563-85-8
     RL: POF (Polymer in formulation); PRP (Properties); TEM (Technical or
     engineered material use); USES (Uses)
        (nanoparticles filled polysilsesquioxanes for to chemical sensing)
IT
     936563-85-8
     RL: POF (Polymer in formulation); PRP (Properties); TEM (Technical or
     engineered material use); USES (Uses)
        (nanoparticle's filled polysilsesquioxanes for to chemical sensing)
RN
     936563-85-8 HCAPLUS
CN
     2-Propenoic acid, 2-methyl-, butyl ester, polymer with
     3-[3,5,7,9,11,13,15-heptakis(2-methylpropyl)pentacyclo[9.5.1.13,9.15,15.17
     ,13]octasiloxan-1-yl]propyl 2-methyl-2-propenoate (CA INDEX NAME)
     CM
          1
     CRN
         307531-94-8
     CMF
         C35 H74 O14 S18
```



CM 2

CRN 97-88-1 CMF C8 H14 O2

$$\begin{array}{c|c} & \text{O} & \text{CH}_2 \\ || & || \\ \text{n-BuO-C-C-Me} \end{array}$$

RE.CNT 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L36 ANSWER 2 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2007:273459 HCAPLUS

DN 146:482346

TI Plasma Surface Modification and Characterization of POSS-Based Nanocomposite Polymeric Thin Films

AU Augustine, Brian H.; Hughes, Wm. Christopher; Zimmermann, Kathryn J.; Figueiredo, Ashley J.; Guo, Xiaowen; Chusuei, Charles C.; Maidment, Jessica S.

CS Department of Chemistry MSC 4501 and Department of Physics, James Madison University, Harrisonburg, VA, 22807, USA

applicant

SO Langmuir (2007), 23(8), 4346-4350 CODEN: LANGD5; ISSN: 0743-7463

PB American Chemical Society

DT Journal

LA English

AB The effect of a remote oxygen plasma on nanocomposite hybrid polymer thin films of poly[(propylmethacryl-heptaisobutyl-polyhedral oligomeric silsesquioxane)-co-(Me methacrylate)] (POSS-MA) has been examined by advancing contact angle, XPS, and variable-angle spectroscopic ellipsometry (VASE). Exposure to a 25 W remote oxygen-containing plasma was found to convert the surface of POSS-MA films from hydrophobic to

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RN

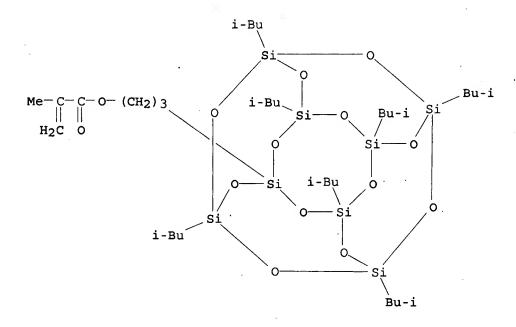
CN

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hydrophilic within 20 s. The exposure time needed for this conversion to
occur decreased as the O2/N2 ratio in the plasma environment increased,
indicating a pos. correlation between the hydrophilicity and the presence
of oxygen in the plasma. Local bonding information inferred from
high-resolution XPS data showed that the iso-Bu bonding to the POSS moiety is
replaced with oxygen as a result of plasma exposure. Finally, VASE data
demonstrates that increasing the weight percent of POSS in the copolymer
significantly impedes the oxygen plasma degradation of POSS-MA films. On the
basis of these results, a model is presented in which the oxygen plasma
removes iso-Bu groups from the POSS cages and leaves a SiO2-like surface
that is correspondingly more hydrophilic than the surface of the untreated
samples and is more resistant to oxidation by the plasma. The ability to
modify surfaces in this manner may impact the utility of this material for
biomedical applications such as microfluidic devices in which
the ability to control surface chemical is critical
35-8 (Chemistry of Synthetic High Polymers)
Section cross-reference(s): 36
silsesquioxane deriv methyl methacrylate copolymer surface property plasma
Thickness
   (film; plasma surface modification and characterization of POSS-based
   nanocomposite polymeric thin films).
Silsesquioxanes
RL: PEP (Physical, engineering or chemical process); PRP (Properties); SPN
(Synthetic preparation); PREP (Preparation); PROC (Process)
   (methacrylate-; plasma surface modification and characterization of
   POSS-based nanocomposite polymeric thin films)
Contact angle
Films
Hybrid organic-inorganic materials
Plasma
Surface composition
Surface treatment
X-ray photoelectron spectroscopy
   (plasma surface modification and characterization of POSS-based
   nanocomposite polymeric thin films)
   (spin; plasma surface modification and characterization of POSS-based
   nanocomposite polymeric thin films)
7727-37-9, Nitrogen, uses
                            7782-44-7, Oxygen, uses
RL: NUU (Other use, unclassified); USES (Uses)
   (plasma surface modification and characterization of POSS-based
   nanocomposite polymeric thin films)
425409-07-0P
RL: PEP (Physical, engineering or chemical process); PRP (Properties); SPN
(Synthetic preparation); PREP (Preparation); PROC (Process)
   (plasma surface modification and characterization of POSS-based
   nanocomposite polymeric thin films)
RL: PEP (Physical, engineering or chemical process); PRP (Properties); SPN
(Synthetic preparation); PREP (Preparation); PROC (Process)
   (plasma surface modification and characterization of POSS-based
  nanocomposite polymeric thin films)
425409-07-0 HCAPLUS
2-Propenoic acid, 2-methyl-, 3-[3,5,7,9,11,13,15-heptakis(2-
methylpropyl)pentacyclo[9.5.1.13,9.15,15.17,13]octasiloxan-1-yl]propyl
ester, polymer with methyl 2-methyl-2-propenoate (CA INDEX NAME)
```

CM 1

CRN 307531-94-8

CMF C35 H74 O14 Si8



CM 2

CRN 80-62-6 CMF C5 H8 O2

RE.CNT 34 THERE ARE 34 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L36 ANSWER 3 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2006:1268071 'HCAPLUS

DN 146:163221

Understanding the reactivity of [W:NAr(CH2tBu)2(:CHtBu)] (Ar =
2,6-iPrC6H3) with silica partially dehydroxylated at low temperatures
through a combined use of molecular and surface organometallic chemistry

AU Rhers, Bouchra; Quadrelli, Elsje Alessandra; Baudouin, Anne; Taoufik, Mostafa; Coperet, Christophe; Lefebvre, Frederic; Basset, Jean-Marie; Fenet, Bernard; Sinha, Amritanshu; Schrock, Richard R.

CS Laboratoire de Chimie Organometallique de Surface (UMR 9986 CNRS/ENSCPE Lyon) 43, Villeurbanne, F-69616, Fr.

SO Journal of Organometallic Chemistry (2006), 691(24-25), 5448-5455 CODEN: JORCAI; ISSN: 0022-328X

PB Elsevier Ltd.

DT Journal

LA English

AB Reaction of [W:NAr(CH2tBu)2(:CHtBu)] (Ar = 2,6-iPrC6H3) with silica partially dehydoxylated at 200° does not lead only to the expected bisgrafted [(.tplbond.SiO)2W:NAr(:CHtBu)] species, but also surface

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RN CN

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reaction intermediates such as [(.tplbond.SiO)2W:NAr(CH2tBu)2]. All these
species were characterized by IR spectroscopy, 1D and 2D solid state NMR,
elemental anal. and mol. models obtained by using silsesquioxanes. While
a mixture of several surface species, the resulting material displays high
activity in the olefin metathesis.
29-11 (Organometallic and Organometalloidal Compounds)
Section cross-reference(s): 66
tungsten imine alkylidene surface reaction partially
dehydoxylated silica NMR
IR spectra
Molecular modeling
Surface reaction
   (NMR study of understanding the reactivity of imine tungsten
   alkylidene with partially dehydoxylated silica at low temps. through
   combined of mol. and surface organometallic chemical)
Silsesquioxanes
RL: RCT (Reactant); RACT (Reactant or reagent)
   (NMR study of understanding the reactivity of imine tungsten
   alkylidene with partially dehydoxylated silica at low temps. through
   combined of mol. and surface organometallic chemical)
Alkenes, reactions
RL: RCT (Reactant); RACT (Reactant or reagent)
   (metathesis model; NMR study of understanding the reactivity of imine
   tungsten alkylidene with partially dehydoxylated silica at low
   temps. through combined of mol. and surface organometallic chemical)
Metathesis
   (olefin; model; NMR study of understanding the reactivity of imine
   tungsten alkylidene with partially dehydoxylated silica at low
   temps. through combined of mol. and surface organometallic chemical)
NMR (nuclear magnetic resonance)
   (solid-state; NMR study of understanding the reactivity of imine
   tungsten alkylidene with partially dehydoxylated silica at low
   temps. through combined of mol. and surface organometallic chemical)
7631-86-9D, Silica, partially dehydroxylated 128951-56-4
329897-28-1
              905703-18-6
RL: RCT (Reactant); RACT (Reactant or reagent)
   (NMR study of understanding the reactivity of imine tungsten
   alkylidene with partially dehydoxylated silica at low temps. through
   combined of mol. and surface organometallic chemical)
919991-99-4P
               919992-01-1P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP
(Preparation); RACT (Reactant or reagent)
   (NMR study of understanding the reactivity of imine tungsten
   alkylidene with partially dehydoxylated silica at low temps. through
   combined of mol. and surface organometallic chemical)
128951-56-4DP, silica modified
                                 905703-17-5P
                                                919992-03-3P
RL: SPN (Synthetic preparation); PREP (Preparation)
   (NMR study of understanding the reactivity of imine tungsten
   alkylidene with partially dehydoxylated silica at low temps. through
   combined of mol. and surface organometallic chemical)
919991-99-4P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP
(Preparation); RACT (Reactant or reagent)
   (NMR study of understanding the reactivity of imine tungsten
   alkylidene with partially dehydoxylated silica at low temps. through
   combined of mol. and surface organometallic chemical)
919991-99-4 HCAPLUS
Tungsten, [2,6-bis(1-methylethyl)benzenaminato(2-)]bis(2,2-
dimethylpropyl)bis(3,5,7,9,11,13,15-heptacyclopentylpentacyclo[9.5.1.13,9.
```

15,15.17,13]octasiloxan-1-olato-κO1)-, (SP-5-21)- (CA INDEX NAME)

PAGE 2-A

RE.CNT 21 THERE ARE 21 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L36 ANSWER 4 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2006:879292 HCAPLUS

DN 146:523461

TI Development of pressure sensitive paints based on silicon nanostructured powders

AU Castaldo, A.; Massera, E.; Quercia, L.; Di Francia, G.

CS Centro Ricerche di Portici, ENEA, 80055 NA, Italy

SO Sensors and Actuators, B: Chemical (2006), B118(1-2), 328-332 CODEN: SABCEB; ISSN: 0925-4005

PB Elsevier B.V.

DT Journal

LA English

AB A new pressure sensitive paint (PSP) family, based on silicon nanostructured powders as luminophore, has been developed for application in stationary wind tunnel test and potential advanced turbomachinery purposes. The pressure sensitive paint operates on its photoluminescence quenching when the nanostructured powder is exposed to an oxygen rich environment. Nanostructured powders, obtained with a combined ball milling-chemical etching process, are synthesized and deposited onto an appropriate polysilsesquioxane matrix, so that our paint can be adjusted for different pressure measurement applications. The formulation we propose is the first example of a PSP composite in which the filler acts as sensible element and it is localized over the polymeric film, reducing the response time due to oxygen variation to only a few seconds (0.5 ÷ 5 s).

CC 42-10 (Coatings, Inks, and Related Products)

ST silicon nanostructured powder pressure sensitive paint

IT Luminescence

Luminescence quenching

Paints

Polymer morphology

(pressure sensitive paints based on silicon nanostructured powders)

IT Silsesquioxanes

RL: PRP (Properties); TEM (Technical or engineered material use); USES (Uses)

(pressure sensitive paints based on silicon nanostructured powders) IT 936563-85-8

RL: PRP (Properties); TEM (Technical or engineered material use); USES (Uses)

(pressure sensitive paints based on silicon nanostructured powders) 936563-85-8

RL: PRP (Properties); TEM (Technical or engineered material use); USES (Uses)

(pressure sensitive paints based on silicon nanostructured powders)

RN 936563-85-8 HCAPLUS

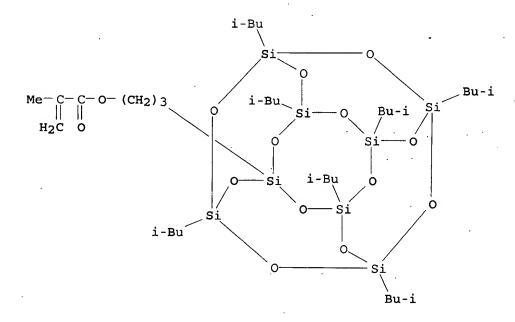
2-Propenoic acid, 2-methyl-, butyl ester, polymer with 3-[3,5,7,9,11,13,15-heptakis(2-methylpropyl)pentacyclo[9.5.1.13,9.15,15.17,13]octasiloxan-1-yl]propyl 2-methyl-2-propenoate (CA INDEX NAME)

CM 1

IT

CN

CRN 307531-94-8 CMF C35 H74 O14 Si8



CM 2

CRN 97-88-1 CMF C8 H14 O2

RE.CNT 17 THERE ARE 17 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

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PENG 10/520763 07/11/2007Page 12
L36
     ANSWER 5 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN
AN
     2006:617113 HCAPLUS
DN
     146:296543
ΤI
     Ultra-low-.vepsiln. nanocomposites of novel fluorinated polyimides grafted
     with polyhedral oligomeric silsesquioxane
ΑU
     Wang, Xiao-Feng; Chen, Yi-Wang
CS
     School of Materials Science and Engineering, Nanchang University,
     Nanchang, 330047, Peop. Rep. China
     Yingyong Huaxue (2006), 23(5), 484-488
SO
     CODEN: YIHUED; ISSN: 1000-0518
PB
     Kexue Chubanshe
DT
     Journal
LA
     Chinese
     Nanocomposites of 6F-Durene with covalently grafted polymethacrylate(PMA)
AΒ
     side chains containing polyhedral oligomeric silsesquioxane (R7R'Si8012 or
     POSS) units were prepared by means of the thermally-initiated free-radical
     graft polymerization of methacrylcyclopentyl-POSS (MA-POSS) with ozone-
     pretreated 6F-Durene. The chemical composition and the structure of the
6F-Durene
     with grafted methacrylcyclopentyl-POSS side chains (POSS/6F-Durene
     copolymers) were characterized by NMR, x-ray diffraction (XRD), and
     field-emission SEM (FESEM). The POSS/6F-Durene nanocomposite films had
     both lower and tunable dielec. consts. (.vepsiln.'s) in
     comparison with the pristine 6F-Durene films, with values between 2.0 and
     2.5.
CC
     37-3 (Plastics Manufacture and Processing)
     Section cross-reference(s): 76
ST
     fluorinated polyimides grafted polyhedral oligomeric silsesquioxane
IT
     Polyimides, preparation
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (fluorine-containing, reaction products; ultra-low-.vepsiln. nanocomposites
        of fluorinated polyimides grafted with polyhedral oligomeric
        silsesquioxane)
IT
     Fluoropolymers, preparation
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (polyimide-, reaction products; ultra-low-.vepsiln. nanocomposites of
        fluorinated polyimides grafted with polyhedral oligomeric
        silsesquioxane)
     Dielectric constant
TT
     Glass transition temperature
     Polymer morphology
        (ultra-low-.vepsiln. nanocomposites of fluorinated polyimides grafted
        with polyhedral oligomeric silsesquioxane)
IT
     928028-21-1P
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (ultra-low-.vepsiln. nanocomposites of fluorinated polyimides grafted
        with polyhedral oligomeric silsesquioxane)
IT
     928028-21-1P
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (ultra-low-.vepsiln. nanocomposites of fluorinated polyimides grafted
        with polyhedral oligomeric silsesquioxane)
RN
     928028-21-1 HCAPLUS
CN
     2-Propenoic acid, 2-methyl-, 3-(3,5,7,9,11,13,15-
     heptacyclopentylpentacyclo[9.5.1.13,9.15,15.17,13]octasiloxan-1-yl)propyl
     ester, polymer with 2,3,5,6-tetramethyl-1,4-benzenediamine and
     5,5'-[2,2,2-trifluoro-1-(trifluoromethyl)ethylidene]bis[1,3-
     isobenzofurandione], graft (CA INDEX NAME)
```

CM

CRN 169391-91-7 CMF C42 H74 O14 Si8

CM 2

CRN 3102-87-2 CMF C10 H16 N2

CM 3

CRN 1107-00-2 CMF C19 H6 F6 O6

```
ANSWER 6 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN
L36
     2006:407280 HCAPLUS
AΝ
DN
     145:83954
TI
     Enhanced efficiency of polyfluorene derivatives: organic-inorganic
     hybrid polymer light-emitting diodes
AU
     Lee, Jonghee; Cho, Hoon-Je; Cho, Nam Sung; Hwang, Do-Hoon; Kang, Jong-Min;
     Lim, Eunhee; Lee, Jeong-Ik; Shim, Hong-Ku
CS
     Center for Advanced Functional Polymers, Department of Chemistry and
     School of Molecular Science (BK21), Korea Advanced Institute of Science
     and Technology, Daejeon, 305-701, S. Korea
SO
     Journal of Polymer Science, Part A: Polymer Chemistry (2006),
     2943-2954
     CODEN: JPACEC; ISSN: 0887-624X
PB
     John Wiley & Sons, Inc.
DT
     Journal
LA
     English
AB
     Two novel organic-inorg. hybrid polyfluorene derivs.,
     poly{(9,9'-dioctyl-2,7-fluorene)-co-(9,9'-di-POSS-2,7-fluorene)-co-[2,5-
     bis(octyloxy)-1,4-phenylene] (PFDOPPOSS) and poly{(9,9'-dioctyl-2,7-
     fluorene)-co-(9,9'-di-POSS-2,7-fluorene)-co-bithiophene} (PFT2POSS), were
     synthesized by the Pd-catalyzed Suzuki reaction of polyhedral oligomeric
     silsesquioxane (POSS) appended fluorene, dioctyl phenylene, and
     bithiophene moieties. The synthesized polymers were characterized with 1H
     NMR spectroscopy and elemental anal. Photoluminescence (PL) studies
     showed that the incorporation of the POSS pendant into the polyfluorene
     derivs. significantly enhanced the fluorescence quantum yields of the
     polymer films, likely via a reduction in the degree of interchain interaction
     as well as keto formation. Addnl., the blue-light-emitting polyfluorene
     derivative PFDOPPOSS showed high thermal color stability in PL. Moreover,
     single-layer light-emitting diode devices of an indium tin
     oxide/poly(3,4-ethylene dioxythiophene):poly(styrene
     sulfonate)/polymer/Ca/Al configuration fabricated with PFDOPPOSS and
     PFT2POSS showed much improved brightness, maximum luminescence intensity, and
     quantum efficiency in comparison with devices fabricated with the
     corresponding pristine polymers PFDOP and PFT2. In particular, the maximum
     external quantum efficiency of PFT2POSS was 0.13%, which was twice that of
     PFT2 (0.06%), and the maximum current efficiency of PFT2POSS was 0.38 cd/A,
     which again was twice that of PFT2 (0.19 cd/A).
CC
     37-3 (Plastics Manufacture and Processing)
     Section cross-reference(s): 36, 73, 76
ST
     octahedral silsesquioxane modification polyfluorene bithiophene
     fluorescence electroluminescence LED
TT
     Band gap
     Electric current-potential relationship
     Electroluminescent devices
     Fluorescent substances
       Hybrid organic-inorganic materials
     Ionization potential
     Luminescence, electroluminescence
     Thermal stability
        (octahedral silsesquioxane-modified polyfluorenes and
        bithiophene-containing polyfluorenes as organic-inorg. hybrid
        polymer light-emitting diodes)
IT
                    457931-26-9P
    239075-02-6P
    RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (monomer; octahedral silsesquioxane-modified polyfluorenes and
        bithiophene-containing polyfluorenes as organic-inorg. hybrid
       polymer light-emitting diodes)
```

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IT
     210347-56-1P
                    850648-80-5P 892142-65-3P
                                                892142-66-4P
     892142-67-5P 892142-68-6P
     RL: PRP (Properties); SPN (Synthetic preparation); PREP
     (Preparation)
        (octahedral silsesquioxane-modified polyfluorenes and
        bithiophene-containing polyfluorenes as organic-inorg. hybrid
        polymer light-emitting diodes)
TT
     111-83-1, 1-Bromooctane 123-31-9, Hydroquinone, reactions
     61676-62-8, 2-Isopropoxy-4,4,5,5-tetramethyl-1,3,2-dioxaborolane
     145483-68-7
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (octahedral silsesquioxane-modified polyfluorenes and
        bithiophene-containing polyfluorenes as organic-inorg. hybrid
        polymer light-emitting diodes)
IT
     67399-94-4P, 1,4-Dioctyloxybenzene
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (octahedral silsesquioxane-modified polyfluorenes and
        bithiophene-containing polyfluorenes as organic-inorg. hybrid
        polymer light-emitting diodes)
IT
     892142-65-3P 892142-68-6P
     RL: PRP (Properties); SPN (Synthetic preparation); PREP
     (Preparation)
        (octahedral silsesquioxane-modified polyfluorenes and
        bithiophene-containing polyfluorenes as organic-inorg. hybrid
        polymer light-emitting diodes)
RN
     892142-65-3 HCAPLUS
     1,3,2-Dioxaborolane, 2,2'-[2,5-bis(octyloxy)-1,4-phenylene]bis[4,4,5,5-
     tetramethyl-, polymer with 2,7-dibromo-9,9-dioctyl-9H-fluorene and
     1,1'-[(2,7-dibromo-9H-fluoren-9-ylidene)bis[2,1-ethanediyloxy-3,1-
     propanediyl (dimethylsilylene) oxy] ] bis [3,5,7,9,11,13,15-
     heptacyclopentylpentacyclo[9.5.1.13,9.15,15.17,13]octasiloxane] (9CI) (CA
     INDEX NAME)
     CM
          1
     CRN 807618-46-8
     CMF C97 H164 Br2 O28 Si18
```

PAGE 1-A

PAGE 2-A

PAGE 4-A

see nept page for Si-O ringe

PAGE 5-A

CM 2

CRN 457931-26-9 CMF C34 H60 B2 O6

CM 3

CRN 198964-46-4 CMF C29 H40 Br2

$$Me-(CH_2)$$
 7 (CH_2) 7 $-Me$ Br

RN 892142-68-6 HCAPLUS

CN 1,3,2-Dioxaborolane, 2,2'-[2,2'-bithiophene]-5,5'-diylbis[4,4,5,5-

KATHLEEN FULLER EIC1700 571/272-2505

tetramethyl-, polymer with 2,7-dibromo-9,9-dioctyl-9H-fluorene and 1,1'-[(2,7-dibromo-9H-fluoren-9-ylidene)bis[2,1-ethanediyloxy-3,1-propanediyl(dimethylsilylene)oxy]]bis[3,5,7,9,11,13,15-heptacyclopentylpentacyclo[9.5.1.13,9.15,15.17,13]octasiloxane] (9CI) (CA INDEX NAME)

CM 1

CRN 807618-46-8 CMF C97 H164 Br2 O28 Si18

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PAGE 2-A

PAGE 3-A

PAGE 4-A

CM 2

CRN 239075-02-6 CMF C20 H28 B2 O4 S2

CM 3

CRN 198964-46-4 CMF C29 H40 Br2

$$Me-(CH_2)$$
 7 (CH_2) 7 $=$ Me Br

RE.CNT 54 THERE ARE 54 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L36 ANSWER 7 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN AN 2006:233054 HCAPLUS

KATHLEEN FULLER EIC1700 571/272-2505

```
DN
     144:451036
     Synthesis, morphology and photophysics of novel hybrid
TT
     organic-inorganic polyhedral oligomeric silsesquioxane-tethered
     poly(fluorenyleneethynylene)s
     Pu, Kan-Yi; Zhang, Bing; Ma, Zhun; Wang, Pei; Qi, Xiao-Ying; Chen,
AU
     Run-Feng; Wang, Lian-Hui; Fan, Qu-Li; Huang, Wei
CS
     Institute of Advanced Materials (IAM), Fudan University, Shanghai, 200433,
     Peop. Rep. China
SO
     Polymer (2006), 47(6), 1970-1978
     CODEN: POLMAG; ISSN: 0032-3861
PB
     Elsevier Ltd.
DT
     Journal
LA
     English
AB
     A series of novel hybrid organic-inorg. light-emitting materials,
     polyhedral oligomeric silsesquioxane-tethered
     poly(fluorenyleneethynylene)s, were successfully synthesized via the
     Sonagashira coupling reaction. The chemical structures of these copolymers
     were determined by 1H NMR and FTIR spectra. The morphologies of these
     copolymers were studied in detail using TEM and WAXD. The WAXD data
     showed that POSS formed small aggregates instead of crystals in the
     polymer matrix, indicating the significant effect of the backbone
     constraint on POSS crystallization Furthermore, it also revealed that the interchain interaction weakened and the interchain distance increased
     after introducing POSS groups. The TEM data indicated that POSS
     aggregates were well dispersed in the polymer matrix. In accordance with
     the morphol. investigation, the results of UV-vis absorption and
     photoluminescence emission spectra of these copolymers showed that the
     tendency toward planar conformation of conjugated backbones was reduced to
     a certain extent due to weakened interchain interaction. Accordingly,
     these copolymers exhibited enhanced quantum yields in the solid state.
     addition, owing to the thermal and oxygen stability of hybrid POSS,
     the thermal spectral stability of these polymers was also improved
     greatly.
CC
     35-7 (Chemistry of Synthetic High Polymers)
ST
     pentacyclooctasiloxane deriv contg polyfluorenyleneethynylene
IT
     Polyacetylenes, preparation
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (cardo, pentacyclooctasiloxane group-containing; synthesis, morphol. and
        photophysics of hybrid organic-inorg. polyhedral oligomeric
        silsesquioxane-tethered poly(fluorenyleneethynylene)s)
IT
     Cardo polymers
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (polyacetylenes, pentacyclooctasiloxane group-containing; synthesis,
        morphol. and photophysics of hybrid organic-inorg. polyhedral
        oligomeric silsesquioxane-tethered poly(fluorenyleneethynylene)s)
IT
    Hybrid organic-inorganic materials
     Luminescence
     Polymer morphology
     UV and visible spectra
        (synthesis, morphol. and photophysics of hybrid organic-inorg.
        polyhedral oligomeric silsesquioxane-tethered
        poly(fluorenyleneethynylene)s)
IT
     14348-75-5, 2,7-Dibromofluorenone
                                          480438-19-5
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (in preparation of monomer for synthesis of hybrid organic-inorg.
        polyhedral oligomeric silsesquioxane-tethered
        poly(fluorenyleneethynylene)s)
IT
     169169-89-5P
    RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
```

(Reactant or reagent)

(in preparation of monomer for synthesis of **hybrid** organic-inorg. polyhedral oligomeric silsesquioxane-tethered poly(fluorenyleneethynylene)s)

IT 885947-40-0P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(monomer; for synthesis hybrid organic-inorg. polyhedral

oligomeric silsesquioxane-tethered poly(fluorenyleneethynylene)s)

IT 885947-43-3P

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)

(synthesis, morphol. and photophysics of hybrid organic-inorg. polyhedral oligomeric silsesquioxane-tethered

poly(fluorenyleneethynylene)s)

IT 885947-40-0P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP
(Preparation); RACT (Reactant or reagent)

(monomer; for synthesis hybrid organic-inorg. polyhedral

oligomeric silsesquioxane-tethered poly(fluorenyleneethynylene)s)

RN 885947-40-0 HCAPLUS

CN Pentacyclo[9.5.1.13,9.15,15.17,13]octasiloxane, 1,1'-[(2,7-dibromo-9H-fluoren-9-ylidene)bis(4,1-phenyleneoxymethylene-4,1-phenylene-2,1-ethanediyl)]bis[heptacyclopentyl- (9CI) (CA INDEX NAME)

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PAGE 2-A

PAGE 3-A

PAGE 4-A

PAGE 5-A

PAGE 5-B

PAGE 6-A

IT 885947-43-3P

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)

(Preparation)

(synthesis, morphol. and photophysics of hybrid organic-inorg. polyhedral oligomeric silsesquioxane-tethered

poly(fluorenyleneethynylene)s)

RN 885947-43-3 HCAPLUS

CN Pentacyclo[9.5.1.13,9.15,15.17,13]octasiloxane, 1,1'-[(2,7-dibromo-9H-fluoren-9-ylidene)bis(4,1-phenyleneoxymethylene-4,1-phenylene-2,1-

ethanediyl)]bis[3,5,7,9,11,13,15-heptacyclopentyl-, polymer with 2,7-dibromo-9-octadecyl-9-octyl-9H-fluorene and 2,7-diethenyl-9-octadecyl-9-octyl-9H-fluorene (9CI) (CA INDEX NAME)

CM 1

CRN 885947-42-2 CMF C43 H66

$$Me^{-(CH_2)}_{17}$$
 $(CH_2)_{7}^{-Me}$ $CH = CH_2$

CM 2

CRN 885947-41-1 CMF C39 H60 Br2

$$Me-(CH_2)_{17}$$
 $(CH_2)_{7}-Me$ Br

CM 3

CRN 885947-40-0 CMF C113 H160 Br2 O26 Si16

PAGE 1-A

PAGE 2-A

PAGE 3-A

PAGE 4-A

PAGE 5-A

PAGE 5-B

PAGE 6-A

R7

RE.CNT 72 THERE ARE 72 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L36 ANSWER 8 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2006:145628 HCAPLUS

DN 144:351846

TI Low- κ nanocomposite films based on polyimides with grafted polyhedral oligomeric silsesquioxane

AU Chen, Yiwang; Chen, Lie; Nie, Huarong; Kang, E. T.

CS School of Materials Science and Engineering, Nanchang University, Nanchang, 330047, Peop. Rep. China

SO Journal of Applied Polymer Science (2006), 99(5), 2226-2232 CODEN: JAPNAB; ISSN: 0021-8995

PB John Wiley & Sons, Inc.

DT Journal

LA English

AB Nanocomposites of polyimides (PI) with covalently grafted polyhedral oligomeric silsesquioxane (R7R'Si8012 or POSS) units were prepared by thermally-initiated free-radical graft polymerization of methacrylcyclopentyl-POSS (MA-POSS) with the ozone-pretreated poly[N,N'-(1,4-phenylene)-3,3',4,4'-benzophenonetetracarboxylic amic acid] (PAA), followed by thermal imidization. The chemical composition and structure of the PI with grafted methacrylcyclopentyl-POSS side chains (PI-g-PMA-POSS copolymers) were characterized by NMR, x-ray diffraction (XRD), and thermogravimetric anal. (TGA). The POSS mols. in each grafted PMA side chain of the amorphous PI films retained the nanoporous crystalline structure, and formed an aggregate of crystallites. The PI-g-PMA-POSS nanocomposite films had both lower and tunable dielec. consts., in comparison with that of the pristine PI films. Dielec. consts. (κ 's) of about 3.0-2.2 were obtained. The present approach offers a convenient way for preparing low-k materials based on existing PI's.

CC 38-3 (Plastics Fabrication and Uses)
Section cross-reference(s): 76

ST low dielec nanocomposite film polyimide grafted polyhedral oligomeric silsesquioxane

IT Polymer morphology

(fracture-surface; low-κ nanocomposite films based on polyimides

```
with grafted polyhedral oligomeric silsesquioxane)
     Density
IT
     Dielectric constant
     Electric insulators
     Glass transition temperature
     Nanocomposites
     Plastic films
     Porosity
        (low-κ nanocomposite films based on polyimides with grafted
        polyhedral oligomeric silsesquioxane)
IT
     Silsesquioxanes
     RL: PRP (Properties); RCT (Reactant); SPN (Synthetic preparation); PREP
     (Preparation); RACT (Reactant or reagent)
        (polyamic acid-, acrylic, cardo, graft, intermediates; low-κ
        nanocomposite films based on polyimides with grafted polyhedral
        oligomeric silsesquioxane)
IT
     Silsesquioxanes
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (polyimide-polyketone-, acrylic, cardo, graft; low-κ
        nanocomposite films based on polyimides with grafted polyhedral
        oligomeric silsesquioxane)
IT
     Polyketones
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (polyimide-silsesquioxane-, acrylic, cardo, graft; low-\kappa
        nanocomposite films based on polyimides with grafted polyhedral
        oligomeric silsesquioxane)
IT
     Polyimides, properties
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (polyketone-silsesquioxane-, acrylic, cardo, graft; low-\kappa
        nanocomposite films based on polyimides with grafted polyhedral
        oligomeric silsesquioxane)
IT
     Fracture surface morphology
        (polymeric; low-κ nanocomposite films based on polyimides with.
        grafted polyhedral oligomeric silsesquioxane)
IT
     Polyamic acids
     RL: PRP (Properties); RCT (Reactant); SPN (Synthetic preparation); PREP
     (Preparation); RACT (Reactant or reagent)
        (silsesquioxane-, acrylic, cardo, graft, intermediates; low-κ
        nanocomposite films based on polyimides with grafted polyhedral
        oligomeric silsesquioxane)
IT
     845508-91-0P
     RL: PRP (Properties); RCT (Reactant); SPN (Synthetic preparation); PREP
     (Preparation); RACT (Reactant or reagent)
        (intermediate; low-k nanocomposite films based on polyimides with
        grafted polyhedral oligomeric silsesquioxane)
IT
     845508-91-0DP, imidized
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (low-κ nanocomposite films based on polyimides with grafted
        polyhedral oligomeric silsesquioxane)
IT
     845508-91-0P
     RL: PRP (Properties); RCT (Reactant); SPN (Synthetic preparation); PREP
     (Preparation); RACT (Reactant or reagent)
        (intermediate; low-k nanocomposite films based on polyimides with
        grafted polyhedral oligomeric silsesquioxane)
RN
     845508-91-0 HCAPLUS
     2-Propenoic acid, 2-methyl-, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,1
CN
     5.17,13]octasiloxanyl)propyl ester, polymer with 1,4-benzenediamine and
     5,5'-carbonylbis[1,3-isobenzofurandione], graft (9CI) (CA INDEX NAME)
```

CM 1

CRN 169391-91-7 CMF C42 H74 O14 Si8

CM 2

CRN 2421-28-5 CMF C17 H6 O7

CM 3

CRN 106-50-3 CMF C6 H8 N2

IT 845508-91-0DP, imidized RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT

KATHLEEN FULLER EIC1700 571/272-2505

PENG 10/520763 07/11/2007Page 32

(Reactant or reagent)

(low- κ nanocomposite films based on polyimides with grafted polyhedral oligomeric silsesquioxane)

RN 845508-91-0 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,1 5.17,13]octasiloxanyl)propyl ester, polymer with 1,4-benzenediamine and 5,5'-carbonylbis[1,3-isobenzofurandione], graft (9CI) (CA INDEX NAME)

CM 1

CRN 169391-91-7 CMF C42 H74 O14 Si8

CM 2

CRN 2421-28-5 CMF C17 H6 O7

CM 3

CRN 106-50-3 CMF C6 H8 N2

RE.CNT 38 THERE ARE 38 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L36 ANSWER 9 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2006:37677 HCAPLUS

DN 145:336914

TI Preparation of fluorinated polyimide/POSS nanocomposites with ultra-low dielectric constant by graft copolymerization

AU Chen, Yiwang; Nie, Huarong; Chen, Lie; Kang, En-Tang

CS School of Materials Science and Engineering, Nanchang University, Nanchang, 330047, Peop. Rep. China

SO Gaofenzi Xuebao (2005), (6), 807-812 CODEN: GAXUE9; ISSN: 1000-3304

PB Kexue Chubanshe

DT Journal

LA Chinese

AB Nanocomposites of fluorinated polyimides (FPI) with covalently grafted polymethacrylate side chains containing polyhedral oligomeric silsesquioxane (R7R'Si8012 or POSS) units were prepared by thermally-initiated free-radical graft polymerization of methacrylcyclopentyl-POSS (MA-POSS) with the ozone-pretreated FPI. The chemical composition and structure of the FPI with grafted methacrylcyclopentyl-POSS side chains were characterized by NMR (1H-NMR) and X-ray diffraction (XRD). The morphol. of the POSS/FPI nanocomposite films was observed by field-emission SEM (FESEM) and transmission electron microscopy (TEM). A layer-by-layer structure was revealed, which supports the formation of an ordered architecture by POSS crystallites in the FPI matrix, as the result of self-assembled POSS units of MA-POSS side chains. The POSS/FPI nanocomposite films had both lower and tunable dielec. consts., in comparison with that of the pristine FPI films. Dielec. consts. (k's) of about 2.5 to 2.1 were obtained. The reduction of dielec. constant was most likely due to a combined contribution of the nanoporosity of the POSS units and the external porosity introduced by the grafting of MA-POSS to the FPI chains.

CC 38-3 (Plastics Fabrication and Uses)

ST fluorinated polyimide POSS nanocomposite dielec const porosity glass temp

IT Polysulfones, uses

RL: POF (Polymer in formulation); PRP (Properties); USES (Uses) (polyether-polyimide-, fluorine-containing; preparation, d., porosity, and dielec. and thermal properties of fluorinated polyimide/POSS nanocomposites with ultra-low dielec. constant)

IT Fluoropolymers, uses

RL: POF (Polymer in formulation); PRP (Properties); USES (Uses) (polyether-polyimide-polysulfone-; preparation, d., porosity, and dielec. and thermal properties of fluorinated polyimide/POSS nanocomposites with ultra-low dielec. constant)

IT Polyimides, uses

RL: POF (Polymer in formulation); PRP (Properties); USES (Uses) (polyether-polysulfone-, fluorine-containing; preparation, d., porosity, and dielec. and thermal properties of fluorinated polyimide/POSS nanocomposites with ultra-low dielec. constant)

IT Polyethers, uses

RL: POF (Polymer in formulation); PRP (Properties); USES (Uses)

PENG 10/520763 07/11/2007Page 34

(polyimide-polysulfone-, fluorine-containing; preparation, d., porosity, and dielec. and thermal properties of fluorinated polyimide/POSS nanocomposites with ultra-low dielec. constant)

IT Density

Dielectric constant

Glass transition temperature

Nanocomposites

Porosity

(preparation, d., porosity, and dielec. and thermal properties of fluorinated polyimide/POSS nanocomposites with ultra-low dielec. constant)

IT 169391-91-7

RL: MOA (Modifier or additive use); PRP (Properties); USES (Uses) (preparation, d., porosity, and dielec. and thermal properties of fluorinated polyimide/POSS nanocomposites with ultra-low dielec. constant)

IT 133028-98-5P 133029-66-0P

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (preparation, d., porosity, and dielec. and thermal properties of fluorinated polyimide/POSS nanocomposites with ultra-low dielec. constant)

IT 169391-91-7

RL: MOA (Modifier or additive use); PRP (Properties); USES (Uses) (preparation, d., porosity, and dielec. and thermal properties of fluorinated polyimide/POSS nanocomposites with ultra-low dielec. constant)

RN 169391-91-7 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,1 5.17,13]octasiloxanyl)propyl ester (9CI) (CA INDEX NAME)

L36 ANSWER 10 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2004:821929 HCAPLUS

DN 142:241025

TI New approach to nanocomposites of polyimides containing polyhedral oligomeric silsesquioxane for dielectric applications

```
ΑU
     Chen, Yiwang; Kang, En-Tang
CS
     School of Material Science and Engineering, Nanchang University, Nanchang,
     330047, Peop. Rep. China
SO
     Materials Letters (2004), 58(29), 3716-3719
     CODEN: MLETDJ; ISSN: 0167-577X
PB
     Elsevier B.V.
DT
     Journal
ĹΑ
     English
AB
     Low dielec. constant nanocomposites of polyimides with grafted methacrylate
     side chains containing polyhedral oligomeric silsesquioxane (POSS) were
     successfully synthesized by thermally initiated free-radical graft
     copolymn. of methacrylcyclopentyl-POSS (MA-POSS) with the
     ozone-preactivated poly(amic acid), followed by thermal imidization.
     dielec. constant of the film can be tuned by varying the molar
     ratio of the grafted MA-POSS side chains in the copolymer.
     37-3 (Plastics Manufacture and Processing)
CC
     Section cross-reference(s): 76
ST
     nanocomposites polyimide contg silsesquioxane dielec application
IT
     Polymerization
        (graft, radical; new approach to nanocomposites of polyimides containing
        polyhedral oligomeric silsesquioxane for dielec. applications)
IT
     Nanocomposites
        (new approach to nanocomposites of polyimides containing polyhedral
        oligomeric silsesquioxane for dielec. applications)
IT
     Silsesquioxanes
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (new approach to nanocomposites of polyimides containing polyhedral
        oligomeric silsesquioxane for dielec. applications)
IT
     Polymer morphology
        (of nanocomposites of polyimides containing polyhedral oligomeric
        silsesquioxane)
ΙT
     Polyketones
     RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation);
     USES (Uses)
        (polyamic acid-, as initiator; in preparation of polyimides containing
        polyhedral oligomeric silsesquioxane for dielec. applications)
IT
     Polyamic acids
     RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation);
     USES (Uses)
        (polyketone-, as initiator; in preparation of polyimides containing
polyhedral
        oligomeric silsesquioxane for dielec. applications)
IT
     Imidation
        (thermal; in preparation of polyimides containing polyhedral oligomeric
        silsesquioxane for dielec. applications)
IT
     845508-91-0DP, imidized
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (assumed and actural monomers; new approach to nanocomposites of
        polyimides containing polyhedral oligomeric silsesquioxane for dielec.
        applications)
IT
     25038-83-9DP, 3,3',4,4'-Benzophenonedicarboxylic anhydride-1,4-
     phenylenediamine copolymer, reaction product with ozone
     RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation);
     USES (Uses)
        (assumed monomers, as initiator; in preparation of polyimides containing
        polyhedral oligomeric silsesquioxane for dielec. applications)
ΙT
     10028-15-6DP, Ozone, reaction product with poly[N,N'-(1,4-phenylene)-
     3,3',4,4'-benzophenonetetracarboxylic amic acid]
     RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation);
     USES (Uses)
```

(in preparation of polyimides containing polyhedral oligomeric silsesquioxane

for dielec. applications)

IT 51396-17-9DP, reaction product with ozone

RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)

(initiator; in preparation of polyimides containing polyhedral oligomeric silsesquioxane for dielec. applications)

IT **845508-91-0DP**, imidized

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (assumed and actural monomers; new approach to nanocomposites of polyimides containing polyhedral oligomeric silsesquioxane for dielec. applications)

RN 845508-91-0 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,1 5.17,13]octasiloxanyl)propyl ester, polymer with 1,4-benzenediamine and 5,5'-carbonylbis[1,3-isobenzofurandione], graft (9CI) (CA INDEX NAME)

CM 1

CRN 169391-91-7 CMF C42 H74 O14 Si8

CM 2

CRN 2421-28-5 CMF C17 H6 O7

CRN 106-50-3 CMF C6 H8 N2

RE.CNT 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L36 ANSWER 11 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2004:101211 HCAPLUS

DN 140:146684

TI Preparation and uses of nonionic telechelic polymers incorporating polyhedral oligosilsesquioxane (POSS)

IN Mather, Patrick T.; Kim, Byoung-Suhk; Ge, Qing; Liu, Changdeng

PA University of Connecticut, USA

SO PCT Int. Appl., 35 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 7

PAN.CNI /																
	PATENT	NO.		KIN	D	DATE		1	APPL:	ICAT:	ION I	NO.		D	ATE	
ΡI	WO 2004	011525	-	A1	-	2004	0205	1	WO 2	 003-1	 US22	 898		20	0030	 723
	W :															
		CO, CR,	CU,	CZ,	DE,	DK,	DM;	DZ,	EC,	EE,	ES,	FI,	GB,	GD,	GE,	GH,
		GM, HR,	HU,	ID,	IL,	IN,	IS,	JP,	ΚE,	KG,	KΡ,	KR,	ΚZ,	LC,	LK,	LR,
		LS, LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NO,	NZ,	OM,	PH,
		PL, PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	TJ,	TM,	TN,	TR,	TT,	TZ,
,		UA, UG,	-	•	•		•	•	•							
	RW:	GH, GM,	KE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	ŪĠ,	ZM,	ZW,	AM,	ΑZ,	BY,
		KG, KZ,	MD,	RU,	ТJ,	TM,	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,
		FI, FR,														
		BF, BJ,														
		024098							US 2	003-	6206	44		20	2030.	716
		606				2006			•							
	AU 2003	254106		A1		2004		1	AU 2	003-	2541	06		20	0030'	723
PRAI		-3995991				2002										
		-620644				2003							•			
•		-4885901		P		2003										
	WO 2003	-US22898	3	W		2003	0723									
GI																

AB Amphiphilic telechelics incorporating polyhedral oligosilsesquioxane (POSS) are synthesized by direct urethane linkage between the diol end groups of polyethylene glycol (PEG) homopolymers and the monoisocyanate group of POSS macromers, wherein the telechelic has a formula (I), wherein R is a cyclic hydrocarbon selected from the group of cyclohexyl, cyclopentyl, cyclooctyl, Me, Et, Pr, iso-Pr, Bu, 'iso-Bu, styryl, vinyl, allyl, ethylphenyl or any aryl, group selected from the group of Ph, biphenyl and naphthyl. The hydrophobicity of the amphiphilic telechelics can be varied by using PEG homopolymers of increasing MW, providing for control over mol. architecture by hydrophilic/hydrophobic balance. A method for synthesizing the amphiphilic telechelic comprises reacting PEG and POSS macromer wherein the monoisocyanate groups of two POSS macromer are directly linked between the diol end groups of PEG. The amphiphilic telechelics are useful as surfactants, thickening agents, additives to plastic such as PMMA (Plexiglass), epoxy adhesives for improved properties.

IC ICM C08G065-32 ICS C08G018-32

CC 35-8 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 37, 46, 52 polyoxyethylene isocyanato oligosilsesquioxane reaction product prepn property; isocyanatodimethylsilylcyclohexyl polyhedral oligosilsesquioxane prepn property use; amphiphilic polyether silsesquioxane telechelic;

toughening agent polyether silsesquioxane

IT Surfactants

ST

(amphiphilic, telechelic; uses of nonionic telechelic polymers incorporating polyhedral oligosilsesquioxane (POSS))

IT Silsesquioxanes

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
 (polyether-; uses of nonionic telechelic polymers incorporating
 polyhedral oligosilsesquioxane (POSS))

```
IT
     Polyoxyalkylenes, reactions
     RL: PRP (Properties); RCT (Reactant); RACT (Reactant or reagent)
        (preparation of nonionic telechelic polymers incorporating polyhedral
        oligosilsesquioxane (POSS))
IT
     Memory devices
        (shape; uses of nonionic telechelic polymers incorporating polyhedral
        oligosilsesquioxane (POSS))
IT
     Polyethers, preparation
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (silsesquioxane-; uses of nonionic telechelic polymers incorporating
        polyhedral oligosilsesquioxane (POSS))
ΙT
     Polymers, preparation
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (telechelic; uses of nonionic telechelic polymers incorporating
        polyhedral oligosilsesquioxane (POSS))
ΙT
     Adhesives
     Battery electrolytes
     Impact modifiers
     Polymer electrolytes
     Thickening agents
        (uses of nonionic telechelic polymers incorporating polyhedral
        oligosilsesquioxane (POSS))
IT
     Epoxy resins, uses
     Polysulfones, uses
     Polyurethanes, uses
     RL: POF (Polymer in formulation); TEM (Technical or engineered material
     use); USES (Uses)
        (uses of nonionic telechelic polymers incorporating polyhedral
        oligosilsesquioxane (POSS))
IT
     77-58-7, Dibutyl tin dilaurate
     RL: CAT (Catalyst use); USES (Uses)
        (preparation of nonionic telechelic polymers incorporating polyhedral
        oligosilsesquioxane (POSS))
IT
                                       476168-98-6
     25322-68-3, Polyethylene glycol
     RL: PRP (Properties); RCT (Reactant); RACT (Reactant or reagent)
        (preparation of nonionic telechelic polymers incorporating polyhedral
        oligosilsesquioxane (POSS))
IT
     476168-99-7P
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (preparation of nonionic telechelic polymers incorporating polyhedral
        oligosilsesquioxane (POSS))
IT
     652973-99-4
     RL: PRP (Properties); TEM (Technical or engineered material use); USES
        (preparation of nonionic telechelic polymers incorporating polyhedral
        oligosilsesquioxane (POSS))
IT
     9003-20-7, Polyvinylacetate
                                  9003-42-3, Polyethylmethacrylate
     9003-53-6D, Polystyrene, sulfonated
                                            9011-14-7,
     Polymethylmethacrylate
     RL: POF (Polymer in formulation); TEM (Technical or engineered material
     use); USES (Uses)
        (uses of nonionic telechelic polymers incorporating polyhedral
        oligosilsesquioxane (POSS))
IT
     476168-99-7P
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (preparation of nonionic telechelic polymers incorporating polyhedral
        oligosilsesquioxane (POSS))
RN
     476168-99-7 HCAPLUS
CN
     Poly(oxy-1,2-ethanediyl), \alpha-[[[3-[[(3,5,7,9,11,13,15-
     heptacyclohexylpentacyclo[9.5.1.13,9.15,15.17,13]octasiloxan-1-
```

yl)oxy]dimethylsilyl]propyl]amino]carbonyl]- ω -[[[[3-[[(3,5,7,9,11,13,15-heptacyclohexylpentacyclo[9.5.1.13,9.15,15.17,13]octas iloxan-1-yl)oxy]dimethylsilyl]propyl]amino]carbonyl]oxy] - (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

PAGE 1-C

PAGE 2-A

PAGE 2-B

RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L36 ANSWER 12 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2004:100839 HCAPLUS

DN 140:146682

TI Synthesis of nonionic telechelic polymers incorporating polyhedral oligosilsesquioxane and uses thereof

IN Mather, Patrick T.; Kim, Byoung-Suhk; Ge, Qing; Liu, Changdeng

PA University of Connecticut, USA

SO U.S. Pat. Appl. Publ., 20 pp. CODEN: USXXCO

DT Patent

LA English

FAN.CNT 7

	PATE	NT N	10.			KIN	D	DATE			APPL	ICAT	ION 1	NO.		D	ATE		
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PΙ	US 2	0040	2409	98		A1	•	2004	0205	1	US. 2	003-	6206	44		2	0030	716	
	US 7	0676	506			B2		2006	0627										
	WO 2	0040	1152	25		A1		2004	0205	1	WO 2	003-	US22	898		2	0030	723	
	1	W:	ΑE,	AG,	ΑL,	AM,	ΑT,	AU,	ΑZ,	BA,	BB,	BG,	BR,	BY,	ΒZ,	CA,	CH,	CN,	
								DK,											
			GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KΕ,	KG,	ΚP,	KR,	ΚZ,	LC,	LK,	LR,	
			LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NO,	ΝZ,	OM,	PH,	

KATHLEEN FULLER EIC1700 571/272-2505

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PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ,
             UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
             KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES,
             FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR,
             BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
     AU 2003254106
                          A1
                                20040216
                                             AU 2003-254106
PRAI US 2002-399599P
                          P
                                 20020730
     US 2003-620644
                          Α
                                 20030716
     US 2003-488590P
                          P
                                 20030718
     WO 2003-US22898
                          W
                                 20030723
GI
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Me Me Si NH-CO
$$\left\{O-CH_2-CH_2\right\}O$$
 CO NH-CO $\left\{O-CH_2-CH_2\right\}O$ NH-CO $\left\{O-$

Amphiphilic telechelics incorporating polyhedral oligosilsesquioxane (POSS) is synthesized by direct urethane linkage between the diol end groups of polyethylene glycol (PEG) homopolymers and the monoisocyanate group of POSS macromers. An amphiphilic telechelic incorporating POSS has the following structure I, wherein R is a cyclic hydrocarbon selected from the group of cyclohexyl, cyclopentyl, cyclooctyl, Me, Et, Pr, iso-Pr, Bu, iso-Bu, styryl, vinyl, allyl, methylphenyl or an aryl group selected from the group of Ph, biphenyl, naphthyl. The hydrophobicity of the amphiphilic telechelics can be varied by using PEG homopolymers of increasing MW, providing for control over mol. architecture by hydrophilic/hydrophobic balance. This amphiphilic telechelics can be used as surfactants, thickening agents, additives to plastic such as PMMA (Plexiglass) and epoxy adhesives for improved properties.

Ι

IC ICM C08J003-00

INCL 524268000

CC 35-8 (Chemistry of Synthetic High Polymers)
Section cross-reference(s): 37, 46

ST polyoxyethylene isocyanato oligosilsesquioxane reaction product prepn property; isocyanatodimethylsilylcyclohexyl polyhedral oligosilsesquioxane prepn property use; amphiphilic surfactant polyether silsesquioxane telechelic; toughening agent polyether silsesquioxane

IT Surfactants

(amphiphilic; uses of nonionic telechelic polymers incorporating polyhedral oligosilsesquioxane)

IT Silsesquioxanes

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (polyether-; synthesis of nonionic telechelic polymers incorporating polyhedral oligosilsesquioxane and uses thereof)

IT Memory devices

(shape; uses of nonionic telechelic polymers incorporating polyhedral oligosilsesquioxane)

IT Polyethers, preparation

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (silsesquioxane-; synthesis of nonionic telechelic polymers incorporating polyhedral oligosilsesquioxane and uses thereof)

IT Polyoxyalkylenes, reactions

RL: PRP (Properties); RCT (Reactant); RACT (Reactant or reagent) (synthesis of nonionic telechelic polymers incorporating polyhedral oligosilsesquioxane and uses thereof)

IT Polymers, preparation

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (telechelic; synthesis of nonionic telechelic polymers incorporating polyhedral oligosilsesquioxane and uses thereof)

IT Impact modifiers

Polymer electrolytes

Thickening agents

(uses of nonionic telechelic polymers incorporating polyhedral oligosilsesquioxane)

IT Epoxy resins, uses

Polysulfones, uses

Polyurethanes, uses

RL: POF (Polymer in formulation); TEM (Technical or engineered material use); USES (Uses)

(uses of nonionic telechelic polymers incorporating polyhedral oligosilsesquioxane)

IT 77-58-7, Dibutyl tin dilaurate

RL: CAT (Catalyst use); USES (Uses)

(synthesis of nonionic telechelic polymers incorporating polyhedral oligosilsesquioxane and uses thereof)

IT 25322-68-3, Polyethylene glycol 476168-98-6

RL: PRP (Properties); RCT (Reactant); RACT (Reactant or reagent) (synthesis of nonionic telechelic polymers incorporating polyhedral oligosilsesquioxane and uses thereof)

IT 476168-99-7P

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (synthesis of nonionic telechelic polymers incorporating polyhedral oligosilsesquioxane and uses thereof)

IT 9003-20-7, Polyvinylacetate 9003-42-3, Polyethylmethacrylate 9003-53-6D, Polystyrene, sulfonated 9011-14-7,

Polymethylmethacrylate

RL: POF (Polymer in formulation); TEM (Technical or engineered material use); USES (Uses)

(uses of nonionic telechelic polymers incorporating polyhedral oligosilsesquioxane)

IT 476168-99-7P

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (synthesis of nonionic telechelic polymers incorporating polyhedral oligosilsesquioxane and uses thereof)

RN 476168-99-7 HCAPLUS

CN Poly(oxy-1,2-ethanediyl), α -[[[3-[[(3,5,7,9,11,13,15-

PAGE 1-A

PAGE 1-B

$$-CH_{2}-CH_{2}$$

$$n$$

$$O$$

$$Si$$

$$O$$

$$Si$$

$$O$$

$$O$$

$$Si$$

PAGE 1-C

PAGE 2-A



PAGE 2-B



RE.CNT 23 THERE ARE 23 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L36 ANSWER 13 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2003:262984 HCAPLUS

DN 139:69571

TI ABA triblock copolymers containing polyhedral oligomeric silsesquioxane pendant groups: synthesis and unique properties

AU Pyun, Jeffrey; Matyjaszewski, Krzysztof; Wu, Jian; Kim, Gyeong-Man; Chun, Seung B.; Mather, Patrick T.

CS Center for Macromolecular Engineering, Department of Chemistry, Carnegie Mellon University, Pittsburgh, PA, 15213, USA

SO Polymer (2003), 44(9), 2739-2750 CODEN: POLMAG; ISSN: 0032-3861

PB Elsevier Science Ltd.

DT Journal

LA English

AB The synthesis and characterization of POSS (polyhedral oligomeric silsesquioxane) containing ABA triblock copolymers is reported. The use of atom transfer radical polymerization (ATRP) enabled the preparation of well-defined

model copolymers possessing a rubbery poly(Bu acrylate) (pBA) middle segment and glassy poly(3-(3,5,7,9,11,13,15-heptaisobutyl-pentacyclo[9.5.1.13,9.15,15.17,13]-octasiloxane-1-yl)propyl methacrylate) (p(MA-POSS)) outer segments. By tuning the relative composition and d.p. (DP) of the two segments, phase separated microstructures were formed in thin films of the copolymer. Specifically, dynamic mech. anal. and transmission electron microscopy (TEM) observations reveal that for a small molar ratio of p(MA-POSS)/pBA (DP = 6/481/6) no evidence of microphase separation is evident while a large ratio (10/201/10) reveals strong microphase separation Surprisingly, the microphase-separated material exhibits a

tensile modulus larger than expected (ca. 2 + 108 Pa) for a continuous rubber phase for temps. between a pBA-related Tg and a softening point for the p(MA-POSS)-rich phase. Transmission electron microscopy (TEM) images with selective staining for POSS revealed the formation of a morphol. consisting of pBA cylinders in a continuous p(MA-POSS) phase. Thermal studies have revealed the existence of two clear glass transitions in the microphase-separated system with strong phys. aging evident for annealing temps. near the Tg of the higher Tg phase (p(MA-POSS)). The observed aging is reflected in wide-angle X-ray scattering as the strengthening of a low-angle POSS-dominated scattering peak,

suggesting some level of ordering during phys. aging. The Tg of the POSS-rich phase observed in the microphase separated triblock copolymer was nearly 25° higher than that of a POSS-homopolymer of the same mol. weight, suggesting a strong confinement-based enhancement of Tg in this

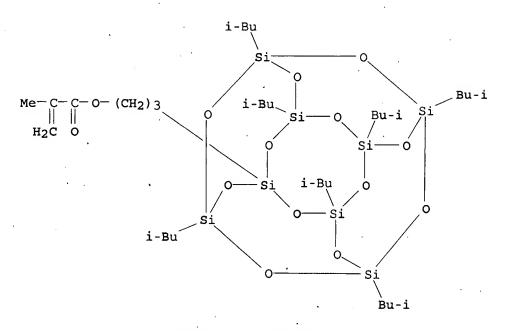
CC 35-4 (Chemistry of Synthetic High Polymers)

ST polyhedral oligomeric silsesquioxane block copolymer prepn thermal mech property

IT · Polymerization

(atom transfer, radical; synthesis and unique properties of ABA triblock copolymers containing polyhedral oligomeric silsesquioxane pendant IT Glass transition temperature Loss modulus Polymer morphology Softening (mechanical) Storage modulus Tensile strength (synthesis and unique properties of ABA triblock copolymers containing polyhedral oligomeric silsesquioxane pendant groups) IT 868-73-5DP, Dimethyl-2,6-dibromoheptanedioate, reaction products with 9003-49-0DP, Poly(butyl acrylate), reaction products poly(Bu acrylate) with dimethyl-2,6-dibromoheptanedioate RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent) (macromer; synthesis and unique properties of ABA triblock copolymers containing polyhedral oligomeric silsesquioxane pendant groups) IT 425409-08-1P RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (synthesis and unique properties of ABA triblock copolymers containing polyhedral oligomeric silsesquioxane pendant groups) IT 255872-36-7P 548798-38-5P 841235-76-5P 842141-07-5P RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (triblock; synthesis and unique properties of ABA triblock copolymers containing polyhedral oligomeric silsesquioxane pendant groups) IT 425409-08-1P RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (synthesis and unique properties of ABA triblock copolymers containing polyhedral oligomeric silsesquioxane pendant groups) ВN 425409-08-1 HCAPLUS CN 2-Propenoic acid, 2-methyl-, 3-[heptakis(2-methylpropyl)pentacyclo[9.5.1.1 3,9.15,15.17,13]octasiloxanyl]propyl ester, homopolymer (9CI) (CA INDEX NAME) CM 1 CRN 307531-94-8

CMF C35 H74 O14 Si8



IT 255872-36-7P 548798-38-5P 841235-76-5P

842141-07-5P

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)

(triblock; synthesis and unique properties of ABA triblock copolymers containing polyhedral oligomeric silsesquioxane pendant groups)

RN 255872-36-7 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,1 5.17,13]octasiloxanyl)propyl ester, polymer with butyl 2-propenoate, block (9CI) (CA INDEX NAME)

CM 1

CRN 169391-91-7 CMF C42 H74 O14 Si8

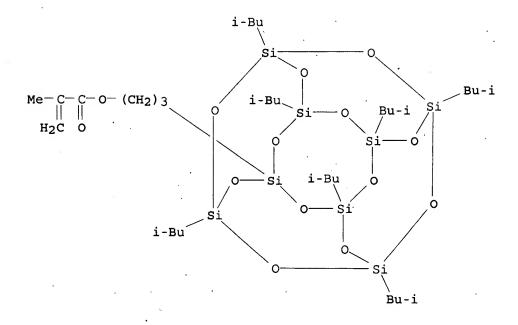
CRN 141-32-2 CMF C7 H12 O2

RN 548798-38-5 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-[heptakis(2-methylpropyl)pentacyclo[9.5.1.1 3,9.15,15.17,13]octasiloxanyl]propyl ester, polymer with butyl 2-propenoate, block (9CI) (CA INDEX NAME)

CM 1

CRN 307531-94-8 CMF C35 H74 O14 Si8



CRN 141-32-2 CMF C7 H12 O2

$$\begin{array}{c} \text{O} \\ \parallel \\ \text{n-BuO-C-CH-----} \text{CH}_2 \end{array}$$

RN 841235-76-5 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,1 5.17,13]octasiloxanyl)propyl ester, polymer with butyl 2-propenoate, triblock (9CI) (CA INDEX NAME)

CM 1

CRN 169391-91-7 CMF C42 H74 O14 Si8

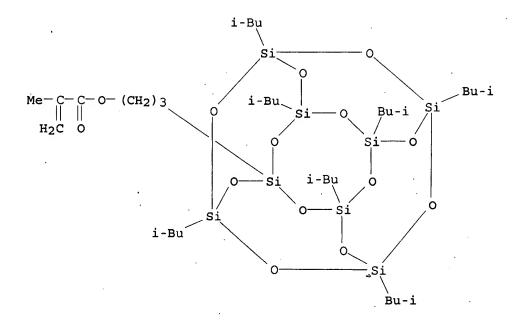
CRN 141-32-2 CMF C7 H12 O2

RN 842141-07-5 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-[heptakis(2-methylpropyl)pentacyclo[9.5.1.1 3,9.15,15.17,13]octasiloxanyl]propyl ester, polymer with butyl 2-propenoate, triblock (9CI) (CA INDEX NAME)

CM 1

CRN 307531-94-8 CMF C35 H74 O14 Si8



CRN 141-32-2 CMF C7 H12 O2

RE.CNT 33 THERE ARE 33 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L36 ANSWER 14 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2000:547502 HCAPLUS

DN 133:135721

TI Functionalizing olefin bearing silsesquioxanes

IN Lichtenhan, Joseph D.; Feher, Frank J.; Soulivong, Daravong

PA United States Dept. of the Air Force, USA

SO U.S., 6 pp.

CODEN: USXXAM

DT Patent

LA English

FAN.CNT 1

				•		
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE	
PI ,	US 6100417	Α	20000808	US 1999-387682	19990831	
PRAT	US 1999-387682		19990831			

AB Reactive silsesquioxanes or polyhedral oligomeric silsesquioxanes (POSS) having ≥1 functionalities, useful as building blocks for hybrid inorg./organic materials, are prepared by cleaving ≥1 olefin bonds in olefin-bearing bearing silsesquioxanes with strong acids or ozone. Thus, Octavinyloctasilsesquioxane ((vinyl)85i8012) is reacted with 1 equiv triflic acid in CH2Cl2 to form TfOCH2CH2(vinyl)7Si8012 at

PENG 10/520763 07/11/2007Page 53

40-45% yield.

IC ICM C07F002-08

INCL 556460000

CC 35-2 (Chemistry of Synthetic High Polymers)

ST silsesquioxane polyhedral oligomeric olefin bearing functionalization; Octavinyloctasilsesquioxane triflate reaction trifluoromethylsulfonyloxyethylheptavinyloctasilsesquioxane prepn

IT Silsesquioxanes

RL: TEM (Technical or engineered material use); USES (Uses) (polyhedral oligomeric; functionalizing olefin bearing silsesquioxanes)

IT . 286389-21-7P

RL: IMF (Industrial manufacture); PREP (Preparation) (functionalizing olefin bearing silsesquioxanes)

(functionalizing olefin bearing silsesquioxanes)

TT 75-75-2, Methanesulfonic acid 1493-13-6, Triflic acid 7664-93-9, Sulfuric acid, reactions 7790-94-5, Chlorosulfuric acid 10028-15-6, Ozone, reactions 69655-76-1, Octavinyloctasilsesquioxane RL: RCT (Reactant); RACT (Reactant or reagent) (functionalizing olefin bearing silsesquioxanes)

IT 286389-21-7P

RL: IMF (Industrial manufacture); PREP (Preparation) (functionalizing olefin bearing silsesquioxanes)

RN 286389-21-7 HCAPLUS

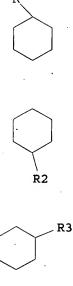
CN Pentacyclo[9.5.1.13,9.15,15.17,13]octasiloxane, 1,1'oxybis[3,5,7,9,11,13,15-heptacyclohexyl- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 2-B



PAGE 3-A



RE.CNT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD

KATHLEEN FULLER EIC1700 571/272-2505

ALL CITATIONS AVAILABLE IN THE RE FORMAT

- L36 ANSWER 15 OF 15 HCAPLUS COPYRIGHT 2007 ACS on STN
- AN 1998:784689 HCAPLUS
- DN 130:139710
- TI Polyhedral oligomeric silsesquioxanes (POSS): silicon based monomers and their use in the preparation of hybrid polyurethanes
- AU Schwab, Joseph J.; Lichtenhan, Joseph D.; Chaffee, Kevin P.; Mather, Patrick T.; Romo-Uribe, Angel
- CS Raytheon STX, Air Force Research Laboratory, Edwards AFB, CA, 93524, USA
- SO Materials Research Society Symposium Proceedings (1998), 519(Organic/Inorganic Hybrid Materials), 21-27 CODEN: MRSPDH; ISSN: 0272-9172
- PB Materials Research Society
- DT Journal
- LA English
- AB A series of polyhedral oligomeric silsesquioxane (POSS) monomers bearing reactive hydroxyl functionalities, suitable for incorporation into step-growth polymers were prepared. These monomers are difunctional in nature and are particularly well suited for use as chain extenders in the synthesis of polyurethanes. Linear segmented polyurethanes prepared using the monomers demonstrated increased modulus, hardness, glass transition temperature, Tg, temperature of 10% weight loss at a heating rate of 10%/min, Tdec, and increased char yields relative to polyurethanes prepared with conventional diol chain extenders. The POSS monomers provided a nanoscale reinforcement of the hard segment domains.
- CC 35-5 (Chemistry of Synthetic High Polymers)
- ST polyhedral oligomeric silsesquioxane chain extender polyurethane
- IT Polyurethanes, preparation
 - RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (block; synthesis of polyhedral oligomeric silsesquioxanes for use as chain extenders in preparation of hybrid block polyurethanes)
- IT 220001-33-2P 220001-34-3P 220001-35-4P
 - RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 - (monomer; for use as chain extenders in preparation of hybrid block polyurethanes)
- IT 682-11-1 1745-89-7, 2,2'-Diallylbisphenol A 220001-32-1
 - RL: RCT (Reactant); RACT (Reactant or reagent)
 - (reactant; in synthesis of polyhedral oligomeric silsesquioxanes for use as chain extenders in preparation of hybrid block polyurethanes)
- IT 220001-36-5P
 - RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
 - (synthesis of polyhedral oligomeric silsesquioxanes for use as chain extenders in preparation of hybrid block polyurethanes)
- IT 220001-34-3P
 - RL: RCT (Reactant); SPN (Synthetic preparation); PREP
 - (Preparation); RACT (Reactant or reagent)
 - (monomer; for use as chain extenders in preparation of hybrid block polyurethanes)
- RN 220001-34-3 HCAPLUS
- CN Phenol, 4,4'-(1-methylethylidene)bis[2-[3-[[(heptacyclohexylpentacyclo[9.5.1.13,9.15,15.17,13]octasiloxanyl)oxy]dimethylsilyl]propyl]- (9CI) (CA INDEX NAME)

PAGE 1-B

PAGE 2-B

IT 220001-36-5P

RL: PRP (Properties); SPN (Synthetic preparation); PREP

(Preparation)

(synthesis of polyhedral oligomeric silsesquioxanes for use as chain extenders in preparation of **hybrid** block polyurethanes)

RN 220001-36-5 HCAPLUS

CN Phenol, 4,4'-(1-methylethylidene)bis[3-[3-[[(heptacyclohexylpentacyclo[9.5.1.13,9.15,15.17,13]octasiloxanyl)oxy]dimethylsilyl]propyl]-, polymer with 1,6-diisocyanatohexane and 4,4'-[oxybis(4,1-butanediyloxy)]bis[1-butanol] (9CI) (CA INDEX NAME)

CM 1

CRN 220001-34-3 CMF C109 H192 O28 Si18

PAGE 1-B

CRN 61136-07-0

CMF C16 H34 O5

 $HO-(CH_2)_4-O-(CH_2)_4-O-(CH_2)_4-O-(CH_2)_4-OH$

CM 3

CRN 822-06-0 CMF C8 H12 N2 O2

OCN-(CH₂)₆-NCO

RE.CNT 13 THERE ARE 13 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

PENG 10/520763 07/11/2007Page 60

NODE ATTRIBUTES: DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

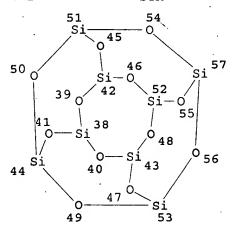
GRAPH ATTRIBUTES: RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 8

STEREO ATTRIBUTES: NONE
L5 9434 SEA FILE=REGISTRY SSS FUL L3
L8 STR

NODE ATTRIBUTES: DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES: RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 5

STEREO ATTRIBUTES: NONE L11 STR



NODE ATTRIBUTES: DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

KATHLEEN FULLER EIC1700 571/272-2505

```
GRAPH ATTRIBUTES:
RING(S) ARE ISOLATED OR EMBEDDED
NUMBER OF NODES IS 20
STEREO ATTRIBUTES: NONE
L19
            361 SEA FILE=REGISTRY SUB=L5 SSS FUL L8 AND L11
L20
            195 SEA FILE=HCAPLUS ABB=ON
                                         L19
              1 SEA FILE=HCAPLUS ABB=ON L20 AND (MICRO(W)FLUID? OR MICROFLUID?
L21
            125 SEA FILE=HCAPLUS ABB=ON L20(L) PREP/RL
L22
L23
              1 SEA FILE=HCAPLUS ABB=ON L22 AND TUNING
L24
              1 SEA FILE=HCAPLUS ABB=ON L19 AND TUNING
L25
              2 SEA FILE=HCAPLUS ABB=ON L21 OR L23 OR L24
L26
             49 SEA FILE=REGISTRY ABB=ON 2 7080.1.1/RID
L28
             41 SEA FILE=HCAPLUS ABB=ON L26
L29
             2 SEA FILE=HCAPLUS ABB=ON L28 AND ?ACRYL?
L30
             32 SEA FILE=HCAPLUS ABB=ON L28(L)PREP/RL
L31
             4 SEA FILE=HCAPLUS ABB=ON L25 OR L29
L32
             28 SEA FILE=HCAPLUS ABB=ON L22 AND HYBRID?
L33
              4 SEA FILE=HCAPLUS ABB=ON L30 AND HYBRID?
L34
             1 SEA FILE=HCAPLUS ABB=ON L30 AND TUN?
L35
              7 SEA FILE=HCAPLUS ABB=ON L20 AND TUN?
L36
             15 SEA FILE=HCAPLUS ABB=ON L31 OR L33 OR L34 OR L35
1.37
             27 SEA FILE=HCAPLUS ABB=ON L32 NOT L36
=> d 137 bib abs ind hitstr 1-27
     ANSWER 1/OF 27 HCAPLUS COPYRIGHT 2007 ACS on STN
L37
AN
     2006:887205 HCAPLUS
DN
     146:501626
TT
     Nanostructured polyurethane-POSS hybrid dispersions by the
     prepolymer mixing process
ΑU
     Nanda, Ajaya K.; Wicks, Douglas A.
     School of Polymers and High Performance Materials, The University of
CS
     Southern Mississippi, Hattiesburg, MS, USA
SO
     PMSE Preprints (2006), 95, 67-69
     CODEN: PPMRA9; ISSN: 1550-6703
PB
     American Chemical Society
DT
     Journal; (computer optical disk)
LA
AΒ
     Waterborne nanostructured polyurethane-POSS (polyhedral oligomeric
     silsesquioxane) hybrid dispersions were synthesized through the
    prepolymer mixing process. To improve the distribution of the POSS
    monomer in the polyurethane aminoethylaminopropylisobutyl polyhedral
    oligomeric silsesquioxane was first pre-reacted with excess isophorone
    diisocyanate in N-methylpyrrolidone. After this reaction the process was
     continued by reaction with poly(hexylene adipate-isophthalate) diol,
     dimethylolpropionic acid, and hexamethylenediamine. Dispersion of the
    polymer was accomplished with tri-Et amine as the neutralizing base. The
    particle size and viscosity of the dispersions were not affected by POSS
    loading. The resulting polyurethane dispersions did show evidence of some
    phase inhomogeneity as evidenced by wide angle X-ray diffraction patterns
```

of cast films. Dynamic mech. anal. showed reinforcement effect reflected

- by the increase in the storage moduli and Tg of the hard segment.
 CC 37-3 (Plastics Manufacture and Processing)
 Section cross-reference(s): 35
- ST polyurethane polyhedral oligomeric silsesquioxane hybrid dispersion mech property

```
10/520763 07/11/2007Page 62
PENG
IT
     Polyurethanes, preparation
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (polyester-polyurea-, block; preparation of nanostructured
        polyurethane-polyhedral oligomeric silsesquioxane hybrid
        dispersions by prepolymer mixing process)
IT
     Polyureas
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (polyester-polyurethane-, block; preparation of nanostructured
        polyurethane-polyhedral oligomeric silsesquioxane hybrid
        dispersions by prepolymer mixing process)
IT
     Polyesters, preparation
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (polyurea-polyurethane-, block; preparation of nanostructured
        polyurethane-polyhedral oligomeric silsesquioxane hybrid
        dispersions by prepolymer mixing process)
IT
     Contact angle
     Disperse systems
     Glass transition temperature
     Loss modulus
     Storage modulus
        (properties of nanostructured polyurethane-polyhedral oligomeric
        silsesquioxane hybrid dispersions prepared by prepolymer mixing
        process)
IT
     Complex modulus
        (tan \delta; properties of nanostructured polyurethane-polyhedral
        oligomeric silsesquioxane hybrid dispersions prepared by
        prepolymer mixing process)
IT
     936448-11-2P 936448-13-4P
     RL: PRP (Properties); SPN (Synthetic preparation); PREP
     (Preparation)
        (composed of actual and assumed monomers; preparation of nanostructured
        polyurethane-polyhedral oligomeric silsesquioxane hybrid
        dispersions by prepolymer mixing process)
IT
     936448-11-2P 936448-13-4P
     RL: PRP (Properties); SPN (Synthetic preparation); PREP
```

(composed of actual and assumed monomers; preparation of nanostructured polyurethane-polyhedral oligomeric silsesquioxane hybrid dispersions by prepolymer mixing process)

RN · 936448-11-2 HCAPLUS

CN 1,3-Benzenedicarboxylic acid, polymer with Desmodur I, N1-[3-[3,5,7,9,11,13,15-heptakis(2-methylpropyl)pentacyclo[9.5.1.13,9.15,1 5.17,13]octasiloxan-1-yl]propyl]-1,2-ethanediamine, hexanedioic acid, 1,6-hexanediol and 3-hydroxy-2-(hydroxymethyl)-2-methylpropanoic acid, block, compd. with N,N-diethylethanamine (CA INDEX NAME)

CM

CRN 121-44-8 CMF C6 H15 N

Εt Et-N-Et

CM

PENG 10/520763 07/11/2007Page 63

CRN 936448-10-1

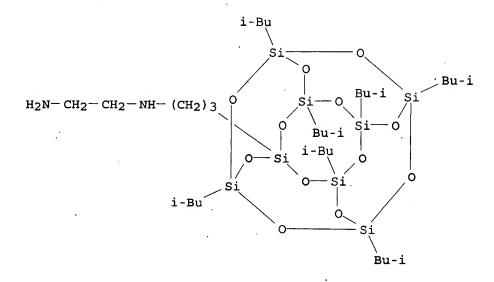
CMF (C33 H76 N2 O12 Si8 . C8 H6 O4 . C6 H14 O2 . C6 H10 O4 . C5 H10 O4 . Unspecified) \mathbf{x}

CCI PMS

CM 3

CRN 444315-16-6

CMF C33 H76 N2 O12 Si8



CM 4

CRN 202149-37-9

CMF Unspecified

CCI PMS, MAN

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

CM 5

CRN 4767-03-7 CMF C5 H10 O4

CM 6

CRN 629-11-8 CMF C6 H14 O2 $HO-(CH_2)_6-OH$

CM 7

CRN 124-04-9 CMF C6 H10 O4

 $HO_2C-(CH_2)_4-CO_2H$

CM 8

CRN 121-91-5 CMF C8 H6 O4

RN 936448-13-4 HCAPLUS

CN 1,3-Benzenedicarboxylic acid, polymer with N1-[3-[3,5,7,9,11,13,15-heptakis(2-methylpropyl)pentacyclo[9.5.1.13,9.15,15.17,13]octasiloxan-1-yl]propyl]-1,2-ethanediamine, hexanedioic acid, 1,6-hexanediol, 3-hydroxy-2-(hydroxymethyl)-2-methylpropanoic acid and 5-isocyanato-1-(isocyanatomethyl)-1,3,3-trimethylcyclohexane, block, compd. with N,N-diethylethanamine (CA INDEX NAME)

CM 1

CRN 121-44-8 CMF C6 H15 N

CM 2

CRN 936448-12-3

CMF (C33 H76 N2 O12 Si8 . C12 H18 N2 O2 . C8 H6 O4 . C6 H14 O2 . C6 H10 O4 . C5 H10 O4) \times

CCI PMS

CM 3

CRN 444315-16-6 CMF C33 H76 N2 O12 Si8

CRN 4767-03-7 CMF C5 H10 O4

CM 5

CRN 4098-71-9 CMF C12 H18 N2 O2

CM 6

CRN 629-11-8 CMF C6 H14 O2

 $^{\rm HO-}$ (CH₂)₆-OH

CRN 124-04-9 CMF C6 H10 O4

 $HO_2C^-(CH_2)_4 - CO_2H$

CM 8

CRN 121-91-5 CMF C8 H6 O4

RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CLYATIONS AVAILABLE IN THE RE FORMAT

L37 ANSWER 2 OF 27 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2006:882317 HCAPLUS

DN 145:455345

TI Nanostructured Polyurethane/POSS Hybrid Aqueous Dispersions Prepared by Homogeneous Solution Polymerization

AU Nanda, Ajaya K.; Wicks, Douglas A.; Madbouly, Samy A.; Otaigbe, Joshua U.

CS School of Polymers and High Performance Materials, University of Southern Mississippi, Hattiesburg, MS, 9406, USA

SO Macromolecules (2006), 39(20), 7037-7043

CODEN: MAMOBX; ISSN: 0024-9297

PB American Chemical Society

DT Journal

LA English

AB Aqueous polyurethane dispersions were prepared with 4-10 wt % of functionalized polyhedral oligomeric silsesquioxanes (POSS) via homogeneous solution polymerization

in acetone followed by solvent exchange with water. The use of acetone as the initial polymerization solvent allowed for the facile incorporation of both diamino and dihydroxy functional POSS monomers in a homogeneous reaction environment. After addition of water and removal of the acetone, stable dispersions with unimodal particle sizes were obtained. The incorporation of the POSS monomers did not have a significant effect on the dispersion's properties; however, the phys. properties of the isolated polymers did display significant changes, with notable increases in storage modulus, Tg, complex viscosity, and surface hydrophobicity. These changes were attributed to the incorporation of the POSS residues into the polyurethane hard segment domains found. Though no sign of any gross phase heterogeneity due to the inclusion of POSS moieties was detected by either thermal characterization or wide-angle X-ray diffraction (WAXD), a significant change was observed by atomic force microscopy (AFM) when the samples were recast from organic solvent.

CC 35-5 (Chemistry of Synthetic High Polymers)
Section cross-reference(s): 36

```
ST
     silsesquioxane contg polyurethane dispersion morphol thermal mech surface
     property
IT
     Polymer morphology
        (domain; nanostructured polyurethane/POSS hybrid aqueous
        dispersions prepared by homogeneous solution polymerization)
     Contact angle
IT
     Dispersion (of materials)
     Elongation at break
     Glass transition temperature
       Hybrid organic-inorganic materials
     Particle size
     Particle size distribution
     Storage modulus
     Tensile strength
     Viscosity
     X-ray diffraction
     Young's modulus
        (nanostructured polyurethane/POSS hybrid aqueous dispersions
        prepared by homogeneous solution polymerization)
IT
     Polyurethanes, preparation
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (polyester-, block, silsesquioxane-substituted; nanostructured
        polyurethane/POSS hybrid aqueous dispersions prepared by
        homogeneous solution polymerization)
IT
     Polyurethanes, preparation
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (polyester-, block; nanostructured polyurethane/POSS hybrid
        aqueous dispersions prepared by homogeneous solution polymerization)
IT
     Polyurethanes, preparation
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (polyester-polyurea-, block, silsesquioxane-substituted; nanostructured
        polyurethane/POSS hybrid aqueous dispersions prepared by
        homogeneous solution polymerization)
IT
     Polyureas
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Préparation)
        (polyester-polyurethane-, block, silsesquioxane-substituted;
        nanostructured polyurethane/POSS hybrid aqueous dispersions
        prepared by homogeneous solution polymerization)
IT
     Polyesters, preparation
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (polyurea-polyurethane-, block, silsesquioxane-substituted;
        nanostructured polyurethane/POSS hybrid aqueous dispersions
        prepared by homogeneous solution polymerization) .
IT
     Softening (mechanical)
        (softening point; nanostructured polyurethane/POSS hybrid aqueous
        dispersions prepared by homogeneous solution polymerization)
IT
     Storage
        (stability; nanostructured polyurethane/POSS hybrid aqueous
        dispersions prepared by homogeneous solution polymerization)
ΙT
     Polymer morphology
        (surface; nanostructured polyurethane/POSS hybrid aqueous
        dispersions prepared by homogeneous solution polymerization)
IT
     Complex modulus
        (tan \delta; nanostructured polyurethane/POSS hybrid aqueous
        dispersions prepared by homogeneous solution polymerization)
     913341-91-0P 913341-93-2P 913341-97-6P
IT
     RL: PRP (Properties); SPN (Synthetic preparation); PREP
     (Preparation)
        (nanostructured polyurethane/POSS hybrid aqueous dispersions
        prepared by homogeneous solution polymerization)
```

IT · 913341-93-2P 913341-97-6P

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)

(nanostructured polyurethane/POSS hybrid aqueous dispersions prepared by homogeneous solution polymerization)

RN 913341-93-2 HCAPLUS

CN 1,3-Benzenedicarboxylic acid, polymer with 1,4-butanediol, Desmodur I, N1-[3-[3,5,7,9,11,13,15-heptakis(2-methylpropyl)pentacyclo[9.5.1.13,9.15,1 5.17,13]octasiloxan-1-yl]propyl]-1,2-ethanediamine, hexanedioic acid, 1,6-hexanediol and 3-hydroxy-2-(hydroxymethyl)-2-methylpropanoic acid, block, compd. with N,N-diethylethanamine (CA INDEX NAME)

CM 1

CRN 121-44-8 CMF C6 H15 N

Et | Et-N-Et

CM 2

CRN 913341-92-1

CMF (C33 H76 N2 O12 Si8 . C8 H6 O4 . C6 H14 O2 . C6 H10 O4 . C5 H10 O4 . C4 H10 O2 . Unspecified) \times

CCI PMS

CM 3

CRN 444315-16-6 CMF C33 H76 N2 O12 Si8

CM 4

PENG 10/520763 07/11/2007Page 69

CRN 202149-37-9 CMF Unspecified CCI PMS, MAN

·

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

CM 5

CRN 4767-03-7 CMF C5 H10 O4

$$\begin{array}{c} \text{Me} & | \\ | \\ \text{HO-CH}_2 - \text{C-CO}_2 \text{H} \\ | \\ \text{CH}_2 - \text{OH} \end{array}$$

CM 6

CRN 629-11-8 CMF C6 H14 O2

$$_{\rm HO^-}$$
 (CH₂)₆-OH

CM 7

CRN 124-04-9 CMF C6 H10 O4

$$HO_2C^-(CH_2)_4^-CO_2H$$

CM · 8

CRN 121-91-5 CMF C8 H6 O4

CM S

CRN 110-63-4 CMF C4 H10 O2 $HO-(CH_2)_4-OH$

RN 913341-97-6 HCAPLUS

CN 1,3-Benzenedicarboxylic acid, polymer with 1,4-butanediol, Desmodur I, 3-[[heptakis(2-methylpropyl)pentacyclo[9.5.1.13,9.15,15.17,13]octasiloxany l]oxy]-1,2-propanediol, hexanedioic acid, 1,6-hexanediol and 3-hydroxy-2-(hydroxymethyl)-2-methylpropanoic acid, block, compd. with N,N-diethylethanamine (9CI) (CA INDEX NAME)

CM 1

CRN 121-44-8 CMF C6 H15 N

Et | Et-N-Et

CM 2

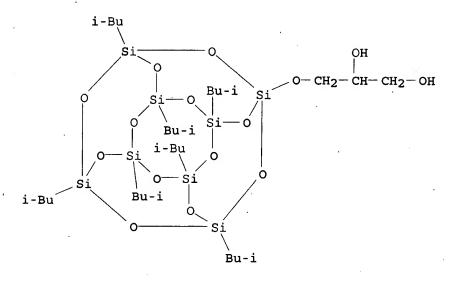
CRN 913341-96-5

CMF (C31 H70 O15 Si8 . C8 H6 O4 . C6 H14 O2 . C6 H10 O4 . C5 H10 O4 . C4 H10 O2 . Unspecified) x

CCI PMS

CM 3

CRN 913341-95-4 CMF C31 H70 O15 Si8



CM 4

CRN 202149-37-9

KATHLEEN FULLER EIC1700 571/272-2505

PENG 10/520763 07/11/2007Page 71

CMF Unspecified CCI PMS, MAN

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

CM 5

CRN 4767-03-7 CMF C5 H10 O4

$$\begin{array}{c} \text{Me} & | \\ | \\ \text{HO-CH}_2 - \text{C-CO}_2 \text{H} \\ | \\ \text{CH}_2 - \text{OH} \end{array}$$

CM 6

CRN 629-11-8 CMF C6 H14 O2

 $_{\rm HO^-}$ (CH₂)₆-OH

CM 7

CRN 124-04-9 CMF C6 H10 O4

 $HO_2C^-(CH_2)_4 - CO_2H$

CM 8

CRN 121-91-5 CMF C8 H6 O4

CM S

CRN 110-63-4 CMF C4 H10 O2

 $HO-(CH_2)_4-OH$

```
RE.CNT 45
              THERE ARE, 45 CITED REFERENCES AVAILABLE FOR THIS RECORD
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
     ANSWER 3 OF 27 HCAPLUS COPYRIGHT 2007 ACS on STN
L37
     2006:827190 HCAPLUS
AN
DN
     145:377630
     Preparation and characterization of polyurethane hybrids from
TТ
     reactive polyhedral oligomeric silsesquioxanes
AU
     Zhang, Shiling; Zou, Qichao; Wu, Limin
CS
     Faculty of Chemistry and Material Science, Hubei University, Wuhan,
     430062, Peop. Rep. China
SO
     Macromolecular Materials and Engineering (2006), 291(7), 895-901
     CODEN: MMENFA; ISSN: 1438-7492
PB
     Wiley-VCH Verlag GmbH & Co. KGaA
DT
     Journal
·LA
     English
AB
     Hybrid polyester resins containing a polyhedral oligomeric
     silsesquioxane (POSS), and their polyurethanes were prepared using a
     reactive POSS (3-[3-[heptakis(2-methylpropyl)pentacyclo[9.5.1.13,9.15,15.1
     7,13]octasil oxanyl]propoxy]-1,2-propanediol) as a substitution for part
     of the diol monomers. They were investigated by FTIR, rheometry, dynamic
     mech. anal., wide-angle x-ray diffraction, contact-angle measurement, atomic
     force microscopy, and thermogravimetric anal., resp. The hybrid
     polyester-POSS resins had a shear thinning behavior, and the more the POSS
     was embedded, the stronger the shear thinning behavior, and the higher the
     viscosity. Incorporation of POSS could increase the glass transition
     temperature and thermal stability and decrease the surface free energy of the
     polyurethanes. When the POSS content was relatively high, the POSS mols.
     in hybrid polyurethane-POSS had a strong self-assembling ability
     to form nanocryst. domains.
     35-5 (Chemistry of Synthetic High Polymers)
CC
st
     polyhedral oligomeric silsesquioxane polyester polyurethane prepn property
IT
     Polyurethanes, preparation
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (polyester-; preparation and characterization of polyester-polyurethane
        hybrids from reactive polyhedral oligomeric silsesquioxane)
IΤ
     Contact angle
     Glass transition temperature
     Surface free energy
     Thermal stability
     Viscosity
        (preparation and characterization of polyester-polyurethane hybrids
        from reactive polyhedral oligomeric silsesquioxane)
IT
     Polyesters, preparation
     RL: CPS (Chemical process); PEP (Physical, engineering or chemical
     process); PRP (Properties); SPN (Synthetic preparation); PREP
     (Preparation); PROC (Process)
        (preparation and characterization of polyester-polyurethane hybrids
        from reactive polyhedral oligomeric silsesquioxane)
IT
     911000-98-1P
     RL: CPS (Chemical process); PEP (Physical, engineering or chemical
     process); SPN (Synthetic preparation); PREP (Preparation); PROC
        (preparation and characterization of polyester-polyurethane hybrids
        from reactive polyhedral oligomeric silsesquioxane)
IT
     911000-99-2P
     RL: PRP (Properties); SPN (Synthetic preparation); PREP
     (Preparation)
        (preparation and characterization of polyester-polyurethane hybrids
```

from reactive polyhedral oligomeric silsesquioxane)

IT 911000-98-1P

RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); SPN (Synthetic preparation); PREP (Preparation); PROC (Process)

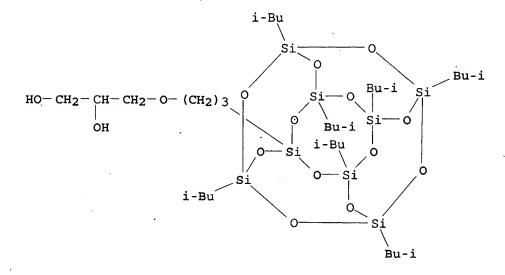
(preparation and characterization of polyester-polyurethane hybrids from reactive polyhedral oligomeric silsesquioxane)

RN 911000-98-1 HCAPLUS

CN Hexanedioic acid, polymer with 1,4-butanediol, 2,2-dimethyl-1,3-propanediol, 3-[3-[heptakis(2-methylpropyl)pentacyclo[9.5.1.13,9.15,15.17, 13]octasiloxanyl]propoxy]-1,2-propanediol and 1,3-isobenzofurandione (9CI) (CA INDEX NAME)

CM 1

CRN 480439-49-4 CMF C34 H76 O15 Si8



CM 2

CRN 126-30-7 CMF C5 H12 O2

CM 3

CRN 124-04-9 CMF C6 H10 O4 HO_2C^- (CH₂)₄ - CO₂H

CM 4

CRN 110-63-4 CMF C4 H10 O2

 $HO-(CH_2)_4-OH$

CM 5

CRN 85-44-9 CMF C8 H4 O3

IT 911000-99-2P

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)

(preparation and characterization of polyester-polyurethane hybrids from reactive polyhedral oligomeric silsesquioxane)

RN 911000-99-2 HCAPLUS

CN Hexanedioic acid, polymer with 1,4-butanediol, 2,2-dimethyl-1,3-propanediol, 3-[3-[heptakis(2-methylpropyl)pentacyclo[9.5.1.13,9.15,15.17, 13]octasiloxanyl]propoxy]-1,2-propanediol, 1,3-isobenzofurandione and 5-isocyanato-1-(isocyanatomethyl)-1,3,3-trimethylcyclohexane (9CI) (CA INDEX NAME)

CM 1

CRN 480439-49-4 CMF C34 H76 O15 Si8

CM 2

CRN 4098-71-9 CMF C12 H18 N2 O2

CM 3

CRN 126-30-7 CMF C5 H12 O2

$$\begin{array}{c} & \text{Me} \\ | \\ \text{HO-} \ \text{CH}_2 - \text{C--} \ \text{CH}_2 - \text{OH} \\ | \\ \text{Me} \end{array}$$

CM 4

CRN 124-04-9 CMF C6 H10 O4

 HO_2C^- (CH₂)₄ - CO_2H

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CM 5

CRN 110-63-4 CMF C4 H10 O2

 $HO-(CH_2)_4-OH$

CM 6

CRN 85-44-9 CMF C8 H4 O3

RE.CNT 34 THERE ARE 34 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 4 OF 27 HCAPLUS COPYRIGHT 2007 ACS on STN L37

AN 2006:550234 HCAPLUS

DN 145:204866

ΤI Lamellar to Inverted Hexagonal Mesophase Transition in DNA Complexes with Calamitic, Discotic, and Cubic Shaped Cationic Lipids

ΑU Cui, Li; Zhu, Lei

CS Polymer Program Institute of Materials Science and Department of Chemical Materials and Biomolecular Engineering, University of Connecticut, Storrs, CT, 06269-3136, USA

SO Langmuir (2006), 22(14), 5982-5985 CODEN: LANGD5; ISSN: 0743-7463

PB American Chemical Society

DT Journal

LΑ English

AΒ In this study, we report on the lipid tail mol. shape/size effect on the mesophase self-assembly behaviors of various cationic lipids complexed with double-stranded DNA. The mol. shape of the cationic lipids was tailored from rodlike (a cyanobiphenyl imidazolium salt) to discotic (a triphenylene imidazolium salt), and finally to cubic [a polyhedral oligomeric silsesquioxane (POSS) imidazolium salt]. An increase in the cross-sectional area of the hydrophobic tails with respect to the hydrophilic imidazolium head induced a neg. spontaneous curvature of the cationic lipids. As a result, a morphol. change from lamello-columnar (LaC) phase for the DNA-cyanobiphenyl imidazolium salt (DNA-rod) and DNA-triphenylene imidazolium salt (DNA-disk) complexes to an inverted hexagonal columnar (HIIC) phase for the DNA-POSS imidazolium salt (DNA-cube) complex was observed The DNA-rod complex had a typical smectic A (SmA) LaC morphol., whereas the DNA-disk complex had a double lamello-columnar liquid crystalline phase. However, when the lipid tail

changed

to POSS, an HIIC morphol. was achieved. These morphol. changes were

CC

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successfully characterized by x-ray diffraction and TEM. We expect that
these liquid crystalline and crystalline DNA hybrid materials may become
potential functional materials for various applications such as organic
microelectronics and gene transfection.
6-7 (General Biochemistry)
Section cross-reference(s): 25, 29, 63, 68, 76
lamellar inverted hexagonal mesophase transition DNA complex cationic
lipid; DNA cyanobiphenyl triphenylene silsesquioxane imidazolium salt
complex mesophase transition; mesophase transition DNA calamitic discotic
cubic lipid cation complex; cyanobiphenyl imidazolium salt prepn DNA
complex mesophase transition; triphenylene imidazolium salt prepn DNA
complex mesophase transition; silsesquioxane imidazolium salt prepn DNA
complex mesophase transition
Lipids, biological studies
RL: BSU (Biological study, unclassified); PRP (Properties); SPN (Synthetic
preparation); BIOL (Biological study); PREP (Preparation)
   (conjugates, with dsDNA; cyanobiphenyl-, silsesquioxane-, and
   triphenylene imidazolium salt cationic lipids preparation and
   lamellar-to-inverted hexagonal mesophase transition self-assembly in
   complexes with dsDNA)
Mesophase
Phase transition
Self-assembly
   (cyanobiphenyl-, silsesquioxane-, and triphenylene imidazolium salt
   cationic lipids preparation and lamellar-to-inverted hexagonal mesophase
   transition self-assembly in complexes with dsDNA)
DNA
RL: BSU (Biological study, unclassified); PRP (Properties); SPN (Synthetic
preparation); BIOL (Biological study); PREP (Preparation)
   (double-stranded, conjugates with cationic lipids; cyanobiphenyl-,
   silsesquioxane-, and triphenylene imidazolium salt cationic lipids
   preparation and lamellar-to-inverted hexagonal mesophase transition
   self-assembly in complexes with dsDNA)
Drug delivery systems
   (liposomes, monolayer (membrane), complexes with DNA; cyanobiphenyl-,
   silsesquioxane-, and triphenylene imidazolium salt cationic lipids
   preparation and lamellar-to-inverted hexagonal mesophase transition
   self-assembly in complexes with dsDNA)
905304-55-4DP, complexes with dsDNA
RL: BSU (Biological study, unclassified); PRP (Properties); SPN (Synthetic
preparation); BIOL (Biological study); PREP (Preparation)
   (calamitic; cyanobiphenyl-, silsesquioxane-, and triphenylene
   imidazolium salt cationic lipids preparation and lamellar-to-inverted
   hexagonal mesophase transition self-assembly in complexes with dsDNA)
905304-57-6DP, complexes with dsDNA
RL: BSU (Biological study, unclassified); PRP (Properties); SPN (Synthetic
preparation); BIOL (Biological study); PREP (Preparation)
   (cubic; cyanobiphenyl-, silsesquioxane-, and triphenylene imidazolium
   salt cationic lipids preparation and lamellar-to-inverted hexagonal
   mesophase transition self-assembly in complexes with dsDNA)
616-47-7, 1-Methylimidazole 15949-84-5, 11-Bromoundecanoyl chloride
19812-93-2
             681235-70-1
RL: RCT (Reactant); RACT (Reactant or reagent)
   (cyanobiphenyl-, silsesquioxane-, and triphenylene imidazolium salt
   cationic lipids preparation and lamellar-to-inverted hexagonal mesophase
   transition self-assembly in complexes with dsDNA)
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RL: RCT (Reactant); SPN (Synthetic preparation); PREP

(cyanobiphenyl-, silsesquioxane-, and triphenylene imidazolium salt

(Preparation); RACT (Reactant or reagent)

83577-85-9P 905304-56-5P

cationic lipids preparation and lamellar-to-inverted hexagonal mesophase transition self-assembly in complexes with dsDNA)

IT 885669-66-9DP, complexes with dsDNA

> RL: BSU (Biological study, unclassified); PRP (Properties); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)

(discotic; cyanobiphenyl-, silsesquioxane-, and triphenylene imidazolium salt cationic lipids preparation and lamellar-to-inverted hexagonal mesophase transition self-assembly in complexes with dsDNA)

IT 905304-57-6DP, complexes with dsDNA

RL: BSU (Biological study, unclassified); PRP (Properties); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)

(cubic; cyanobiphenyl-, silsesquioxane-, and triphenylene imidazolium salt cationic lipids preparation and lamellar-to-inverted hexagonal mesophase transition self-assembly in complexes with dsDNA)

RN 905304-57-6 HCAPLUS

CN 1H-Imidazolium, 1-[11-[3-[heptakis(2-methylpropyl)pentacyclo[9.5.1.13,9.15 ,15.17,13]octasiloxanyl]propoxy]-11-oxoundecyl]-3-methyl-, bromide (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 2-A

● Br⁻

ONE OR MORE TAUTOMERIC DOUBLE BONDS NOT DISPLAYED IN THE STRUCTURE IT

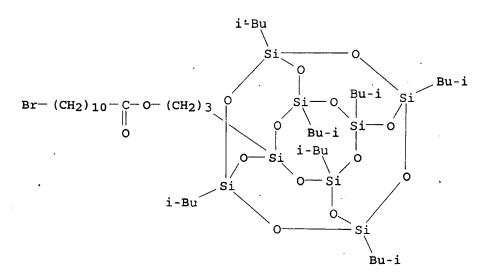
905304-56-5P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(cyanobiphenyl-, silsesquioxane-, and triphenylene imidazolium salt cationic lipids preparation and lamellar-to-inverted hexagonal mesophase transition self-assembly in complexes with dsDNA)

905304-56-5 HCAPLUS RN

Undecanoic acid, 11-bromo-, 3-[heptakis(2-methylpropyl)pentacyclo[9.5.1.13 CN ,9.15,15.17,13]octasiloxanyl]propyl ester (9CI) (CA INDEX NAME)



RE.CNT THERE ARE 24 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

```
10/520763 07/11/2007Page 80
L37
     ANSWER 5 OF 27
                     HCAPLUS COPYRIGHT 2007 ACS on STN
     2006:206761 HCAPLUS
AΝ
DN
     144:451583
     Structure-Property Relationships in Organic-Inorganic Nanomaterials Based
TT
     on Methacryl-POSS and Dimethacrylate Networks
AU
     Bizet, Stephane; Galy, Jocelyne; Gerard, Jean-Francois
CS
     Laboratoire des Materiaux Macromoleculaires/IMP UMR CNRS 5627, Institut
     National des Sciences Appliquees de Lyon, Villeurbanne, F-69621, Fr.
SO
     Macromolecules (2006), 39(7), 2574-2583
     CODEN: MAMOBX; ISSN: 0024-9297
PB
     American Chemical Society
DT
     Journal
LA
     English
AB
     Dimethacrylate-based networks were modified with well-defined organic-inorg.
     building blocks, i.e., polyhedral oligomeric silsesquioxanes (POSS). POSS
     were incorporated in the polymer networks either as dangling chains or as
     crosslinking points. For this purpose, the POSS functionality was varied.
     The influence of the structure of the organic substituent of the POSS cage
     was also investigated. The structure of the POSS-modified networks was
     determined by WAXS, TEM, and thermomech. anal. The POSS as a pendant unit on
     the network backbone shows a strong tendency toward aggregation and
     crystallization, depending on the nature of the organic ligands. The POSS-POSS
     interaction was found to be the main parameter governing the network
     morphol. However, dynamic mech. properties remain nearly at the same
     Tevel as the neat matrix. Multifunctional POSS shows a higher miscibility
     with the dimethacrylate monomer and disperses very well in the cured
     network. As expected, the rubbery modulus grows with increasing amts. of
     POSS according to the high functionality of these addnl. cross-links,
     whereas the glass transition temperature remains constant It is also
demonstrated
     that if the polymerization occurs at high temperature, the distribution of
relaxation
     times is reduced and more homogeneous hybrid networks, in terms
     of mol. mobility, are obtained.
CC
     38-3 (Plastics Fabrication and Uses)
ST
     POSS org inorg nanomaterial structure property relationship;
     silsesquioxane cage polymer org inorg nanomaterial
TT
     Mechanical properties
        (dynamic; structure-property relationships in organic-inorg. nanomaterials
        based on methacryl-POSS and dimethacrylate networks)
ΤT
     Hybrid organic-inorganic materials
     Miscibility
     Nanocomposites
        (structure-property relationships in organic-inorg, nanomaterials based on
        methacryl-POSS and dimethacrylate networks)
IT
     Silsesquioxanes
     RL: POF (Polymer in formulation); PRP (Properties); TEM (Technical or
     engineered material use); USES (Uses)
        (structure-property relationships in organic-inorg. nanomaterials based on
        methacryl-POSS and dimethacrylate networks)
IT
     885479-88-9P 885479-89-0P 885479-90-3P
     RL: PRP (Properties); SPN (Synthetic preparation); TEM (Technical or
     engineered material use); PREP (Preparation); USES (Uses)
        (structure-property relationships in organic-inorg. nanomaterials based on
        methacryl-POSS and dimethacrylate networks)
```

RL: PRP (Properties); SPN (Synthetic preparation); TEM (Technical or

(structure-property relationships in organic-inorg. nanomaterials based on

engineered material use); PREP (Preparation); USES (Uses)

methacryl-POSS and dimethacrylate networks)

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885479-88-9P 885479-89-0P 885479-90-3P

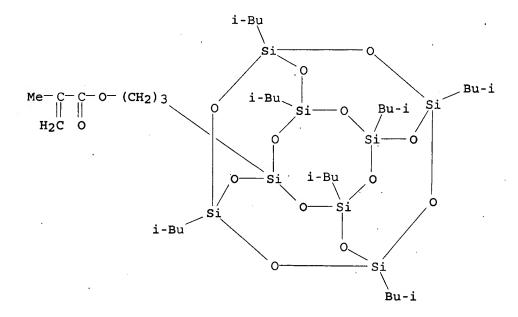
IT

RN 885479-88-9 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, (1-methylethylidene)bis(4,1-phenyleneoxy-2,1-ethanediyloxy-2,1-ethanediyl) ester, polymer with cyclohexyl 2-methyl-2-propenoate and 3-[heptakis(2-methylpropyl)pentacyclo[9.5.1.13,9.15,15.17,13]octasiloxanyl]propyl 2-methyl-2-propenoate (9CI) (CA INDEX NAME)

CM 1

CRN 307531-94-8 CMF C35 H74 O14 Si8



CM 2

CRN 56744-60-6 CMF C31 H40 O8

PAGE 1-A

PAGE 1-B

$$\begin{array}{c|c} & \text{O} & \text{CH}_2 \\ & || & || \\ --\text{CH}_2 - \text{O} - \text{CH}_2 - \text{CH}_2 - \text{O} - \text{C} - \text{C} - \text{Me} \end{array}$$

CM 3

CRN 101-43-9 CMF C10 H16 O2

RN 885479-89-0 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, (1-methylethylidene)bis(4,1-phenyleneoxy-2,1-ethanediyloxy-2,1-ethanediyl) ester, polymer with cyclohexyl 2-methyl-2-propenoate and 3-(heptacyclohexylpentacyclo[9.5.1.13,9.15,15.17,13]octasiloxanyl)propyl 2-methyl-2-propenoate (9CI) (CA INDEX NAME)

CM 1

CRN 169391-90-6 CMF C49 H88 O14 Si8

PENG 10/520763 07/11/2007Page 83

CM 2

CRN 56744-60-6 CMF C31 H40 O8

PAGE 1-A

PAGE 1-B

$$\begin{array}{c|c} & \text{O} & \text{CH}_2 \\ & || & || \\ -\text{CH}_2 - \text{O} - \text{CH}_2 - \text{CH}_2 - \text{O} - \text{C} - \text{C} - \text{Me} \end{array}$$

CM 3

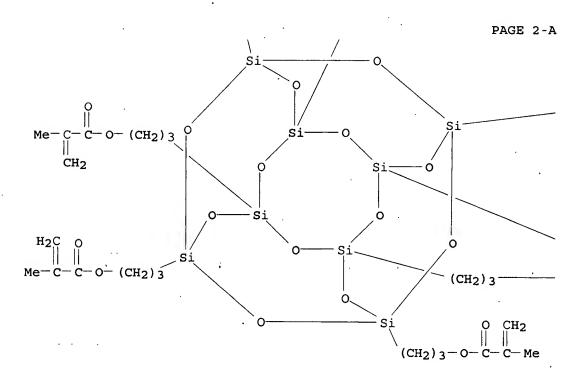
CRN 101-43-9 CMF C10 H16 O2

RN 885479-90-3 HCAPLUS CN 2-Propenoic acid, 2-r

2-Propenoic acid, 2-methyl-, pentacyclo[9.5.1.13,9.15,15.17,13]octasiloxan e-1,3,5,7,9,11,13,15-octaylocta-3,1-propanediyl ester, polymer with (1-methylethylidene)bis(4,1-phenyleneoxy-2,1-ethanediyloxy-2,1-ethanediyl)bis(2-methyl-2-propenoate) (9CI) (CA INDEX NAME)

CM 1

CRN 622404-27-7 CMF C56 H88 O28 Si8



$$\begin{array}{c|c} & \text{O} & \text{CH}_2 \\ & \parallel & \parallel \\ - & \text{(CH}_2)_3 - \text{O} - \text{C} - \text{C} - \text{Me} \end{array}$$

CM 2

CRN 56744-60-6 CMF C31 H40 O8

PAGE 1-A

$$H_{2}^{C}$$
 O $\| \ \| \ \|$ Me-C-C-O-CH₂-CH₂-O-CH₂-CH₂-O $\| \ \| \ \|$ Me

PAGE 1-B

$$\begin{array}{c|c} & \text{O} & \text{CH}_2 \\ & || & || \\ -\text{CH}_2 - \text{O} - \text{CH}_2 - \text{CH}_2 - \text{O} - \text{C} - \text{C} - \text{Me} \end{array}$$

RE.CNT 37 THERE ARE 37 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L37 ANSWER 6 OF 27 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2005:812440 HCAPLUS

DN 143:347228

TI Modification and characterization of Si-based nanobuilding blocks precursors for hybrid materials

AU Mammeri, Fayna; Douja, Najiba; Bonhomme, Christian; Ribot, Francois; Babonneau, Florence; Dire, Sandra

KATHLEEN FULLER EIC1700 571/272-2505

```
Dipartimento di Ingegneria dei Materiali e Tecnologie Industriali,
CS
     Universita di Trento, Trento, 38050, Italy
SO
     Materials Research Society Symposium Proceedings (2005), Volume Date 2004,
     847 (Organic/Inorganic Hybrid Materials--2004), 363-368
     CODEN: MRSPDH; ISSN: 0272-9172
PR
     Materials Research Society
DT
     Journal
LΑ
     English
AB
     New functional nanobuilding blocks were successfully synthesized by
     hydrosilylation of unsatd. alcs. with dimethylsiloxy isobutyl-POSS and
     further acylation with methacryloyl chloride. The solvent influence on
     the reaction pathway was studied, and reaction steps and final
     nano-objects were characterized using multinuclear NMR and FTIR
     spectroscopy. The organic spacer chain length between the inorg. cage and
     the reactive methacrylate function was changed to modify the reactivity of
     the final functional nanobuilding block in the polymerization process.
CC
     29-6 (Organometallic and Organometalloidal Compounds)
ST
     hydrosilylation unsatd alc dimethylsiloxy polyhedral oligomeric
     silsesquioxane Karstedt catalyst
IT
     Hydrosilylation
        (hydrosilylation of unsatd. alcs. with dimethylsiloxy polyhedral
        oligomeric silsesquioxane catalyzed by Karstedt's catalyst)
IT
     Cage compounds
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (hydrosilylation of unsatd. alcs. with dimethylsiloxy polyhedral
        oligomeric silsesquioxane catalyzed by Karstedt's catalyst)
IT
     Acylation
        (hydrosilylation of unsatd. alcs. with dimethylsiloxy polyhedral
        oligomeric silsesquioxane catalyzed by Karstedt's catalyst followed by
IT
     NMR (nuclear magnetic resonance)
        (multinuclear; hydrosilylation of unsatd. alcs. with dimethylsiloxy
        polyhedral oligomeric silsesquioxane catalyzed by Karstedt's catalyst)
ΙT
     Alcohols, reactions
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (unsatd.; hydrosilylation of unsatd. alcs. with dimethylsiloxy
        polyhedral oligomeric silsesquioxane catalyzed by Karstedt's catalyst)
IT
     RL: CAT (Catalyst use); USES (Uses)
        (hydrosilylation of unsatd. alcs. with dimethylsiloxy polyhedral
        oligomeric silsesquioxane catalyzed by Karstedt's catalyst)
     107-18-6, Allyl alcohol, reactions
                                          821-09-0, 4-Penten-1-ol
IT
                                                                     920-46-7,
     Methacryloyl chloride
                             701301-39-5
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (hydrosilylation of unsatd. alcs. with dimethylsiloxy polyhedral
        oligomeric silsesquioxane catalyzed by Karstedt's catalyst)
IT
     866034-88-0P
                    866034-89-1P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent).
        (hydrosilylation of unsatd. alcs. with dimethylsiloxy polyhedral
        oligomeric silsesquioxane catalyzed by Karstedt's catalyst)
IT
                    866034-87-9P 866034-90-4P 866034-91-5P
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (hydrosilylation of unsatd. alcs. with dimethylsiloxy polyhedral
        oligomeric silsesquioxane catalyzed by Karstedt's catalyst)
     866034-90-4P 866034-91-5P
IT
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (hydrosilylation of unsatd. alcs. with dimethylsiloxy polyhedral
        oligomeric silsesquioxane catalyzed by Karstedt's catalyst)
RN
     866034-90-4 HCAPLUS
```

PENG 10/520763 07/11/2007Page 87

CN 2-Propenoic acid, 2-methyl-, 3-[[[heptakis(2-methylpropyl)pentacyclo[9.5.1 .13,9.15,15.17,13]octasiloxanyl]oxy]dimethylsilyl]propyl ester (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 2-A

$$\begin{array}{c} \text{CH}_2 \\ || \\ \text{Me--} \text{C---} \text{R} \end{array}$$

RN 866034-91-5 HCAPLUS
CN 2-Propenoic acid, 2-methyl-, 5-[[[heptakis(2-methylpropyl)pentacyclo[9.5.1 .13,9.15,15.17,13]octasiloxanyl]oxy]dimethylsilyl]pentyl ester (9CI) (CA INDEX NAME)

PAGE 1-A

CH₂

RE.CNT 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CYTATIONS AVAILABLE IN THE RE FORMAT

L37 ANSWER 7 OF 67 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2005:733814 HCAPLUS

DN 144:274608

TI Block copolymer-derived nano-templated surfaces for investigation of nanophysics of wetting

AU Intasanta, Narupol; Coughlin, E. Bryan; Russell, Thomas P.

CS Department of Polymer Science and Engineering, University of Massachusetts Amherst, Amherst, MA, 01003, USA

SO PMSE Preprints (2005), 93, 725-726 CODEN: PPMRA9; ISSN: 1550-6703

PB American Chemical Society

DT Journal; (computer optical disk)

LA English

AB Organic-inorg. hybrid diblock copolymers of a polyhedral oligomeric silsesquioxane methacrylate (MAPOSS) and Me methacrylate (MMA) were prepared by atom transfer radical polymerization. The microphase separation of the block copolymers gave rise to organic-inorg. hybrid nanostructures with different morphologies. The surface-induced orientation of the nanostructures resulted in a nano-templated surface with height modulations corresponding to the domain sizes of the block copolymers. Thermolysis was used to erode the organic components and PMMA, while the inorg. component (POSS) was transformed into a silicon dioxide nanotemplate. Surface modification with various chlorosilanes provided means to study wetting phenomena of the nano-templated surfaces.

CC 35-4 (Chemistry of Synthetic High Polymers)

ST polyhedral oligomeric silsesquioxane methyl methacrylate block copolymer nanostructure; surface modified silica nanotemplate wetting investigation IT Wetting

(block copolymer-derived nano-templated surfaces for investigation of)

IT Templates

(block copolymer-derived nano-templated surfaces for investigation of wetting)

IT Polymer morphology

(surface; block copolymer-derived nano-templated surfaces for investigation of wetting)

TT 7631-86-9P, Silica, preparation 705926-41-6P

RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); SPN (Synthetic preparation); PREP (Preparation); PROC (Process)

(block copolymer-derived nano-templated surfaces for investigation of wetting)

IT 878204-84-3

RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process)

(modifier for nano-templated silica surface; block copolymer-derived nano-templated surfaces for investigation of wetting)

IT 705926-41-6P

RL: CPS (Chemical process); PEP (Physical, engineering or chemical

PENG 10/520763 07/11/2007Page 89

process); SPN (Synthetic preparation); PREP (Preparation); PROC
(Process)

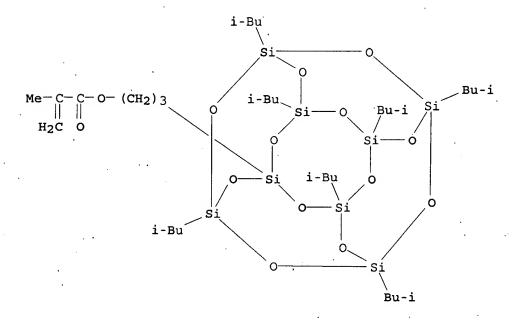
(block copolymer-derived nano-templated surfaces for investigation of wetting)

RN 705926-41-6 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-[heptakis(2-methylpropyl)pentacyclo[9.5.1.1 3,9.15,15.17,13]octasiloxanyl]propyl ester, polymer with methyl 2-methyl-2-propenoate, diblock (9CI) (CA INDEX NAME)

CM 1

CRN 307531-94-8 CMF C35 H74 O14 Si8



CM 2

CRN 80-62-6 CMF C5 H8 O2

$$^{\mathrm{H_2C}}_{\phantom{\mathrm{C}}\phantom{\mathrm{C}}\phantom{\mathrm{O}}\phantom{\mathrm{O}}\phantom{\mathrm{O}}}$$

RE.CNT 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L37 ANSWER 8 OF 27 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2005:229125 HCAPLUS

DN 142:464109

TI Synthesis of the Organic/Inorganic Hybrid Star Polymers and Their Inclusion Complexes with Cyclodextrins

AU Chan, Shih-Chi; Kuo, Shiao-Wei; Chang, Feng-Chih

CS Institute of Applied Chemistry, National Chiao Tung University, Hsin Chu,

851588-81-3P 851588-82-4P

ΙT

RL: RCT (Reactant); RACT (Reactant or reagent)

RL: SPN (Synthetic preparation); PREP (Preparation)

(silylation with (tert-Butyldimethylsilyloxy)propylene)

(preparation and characterization of)

RN851588-81-3 HCAPLUS

CN α -Cyclodextrin, compd. with $\alpha, \alpha', \alpha'', \alpha'''$, a lpha.'''', α''''', α'''''', α''''''-[pentacyclo[9.5.1.13,9.15,15.17,13]octasiloxane-1,3,5,7,9,11,13,15-

octayloctakis[oxy(dimethylsilylene)-3,1-propanediyl]]octakis[ωhydroxypoly[oxy(1-oxo-1,6-hexanediyl)]] (9CI) (CA INDEX NAME)

CM

CRN 851588-80-2

CMF (C6 H10 O2)n C40 H104 O28 Si16 CCI PMS

STRUCTURE DIAGRAM IS NOT AVAILABLE ***

CM

10016-20-3 CRN CMF C36 H60 O30

Absolute stereochemistry.

RN851588-82-4 HCAPLUS

CN γ -Cyclodextrin, compd. with $\alpha, \alpha', \alpha'', \alpha'''$, a lpha.'''', α''''', α'''''', α''''''-[pentacyclo[9.5.1.13,9.15,15.17,13]octasiloxane-1,3,5,7,9,11,13,15octayloctakis [oxy(dimethylsilylene)-3,1-propanediyl]] octakis [ω hydroxypoly[oxy(1-oxo-1,6-hexanediyl)]] (9CI) (CA INDEX NAME)

CM 1

CRN 851588-80-2

CMF (C6 H10 O2)n C40 H104 O28 Si16 CCI **PMS**

STRUCTURE DIAGRAM IS NOT AVAILABLE ***

CM 2

CRN 17465-86-0 C48 H80 O40 CMF

IT 851588-80-2P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP

(Preparation); RACT (Reactant or reagent)

(preparation of organic-inorg. hybrid star polycaprolactones and their inclusion complexes with cyclodextrins)

RN 851588-80-2 HCAPLUS

Poly[oxy(1-oxo-1,6-hexanediyl)], $\alpha,\alpha',\alpha'',\alpha'''$, al CN pha. $(1111, \alpha^{1})(111, \alpha^{1$ [pentacyclo[9.5.1.13,9.15,15.17,13]octasiloxane-1,3,5,7,9,11,13,15octayloctakis[oxy(dimethylsilylene)-3,1-propanediyl]]octakis[ω-

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

hydroxy- (9CI) (CA INDEX NAME)

RE.CNT 56 THERE ARE 56 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L37 ANSWER 9 OF 27 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2005:200878 HCAPLUS

DN

TI Synthesis and cell viability studies of polyhedral oligosilsesquioxane polycaprolactone hybrid materials

AU Skaria, Sunny; Schricker, Scott

CS College of Dentistry, The Ohio State University, Columbus, OH, 43210, USA

Polymer Preprints (American Chemical Society, Division of Polymer SO Chemistry) (2005), 46(1), 94-95 CODEN: ACPPAY; ISSN: 0032-3934

PB American Chemical Society, Division of Polymer Chemistry

RL: PAC (Pharmacological activity); PRP (Properties); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP

polycaprolactone- and polyester-polyhedral oligosilsesquioxane

(multi-arm star; synthesis and cell viability studies of biodegradable

(Preparation); USES (Uses)

hybrid materials)

```
PENG 10/520763 07/11/2007Page 94
RN
     851588-80-2 HCAPLUS
CN
     Poly[oxy(1-oxo-1,6-hexanediyl)], \alpha,\alpha',\alpha'',\alpha''', al
     pha. '''', \alpha''''', \alpha'''''', \alpha''''''-
     [pentacyclo[9.5.1.13,9.15,15.17,13]octasiloxane-1,3,5,7,9,11,13,15-
     octayloctakis[oxy(dimethylsilylene)-3,1-propanediyl]]octakis[ω-
     hydroxy- (9CI)
                    (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
RN
     862667-09-2 HCAPLUS
CN
     Poly[oxy[(1S)-1-methyl-2-oxo-1,2-ethanediyl]],
     ω,ω',ω'',ω''',ω'''',ω'''',ω''''
     '',ω''''-[pentacyclo[9.5.1.13,9.15,15.17,13]octasiloxane-
     1,3,5,7,9,11,13,15-octayloctakis(oxy(dimethylsilylene)-3,1-
     propanediyloxy]]octakis[\alpha-hydro- (9CI) (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
RE.CNT 8
              THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
     ANSWER 10 OF 27/ HCAPLUS COPYRIGHT 2007 ACS on STN
AN
     2005:106619 HCAPLUS
DN
     142:355659
TI
     Organic-Inorganic Poly (Methyl Methacrylate) Hybrids with
     Confined Polyhedral Oligosilsesquioxane Macromonomers
     Toepfer, Oliver; Neumann, Daniel; Choudhury, Namita Roy; Whittaker,
AU
     Andrew; Matisons, Janis
CS
     Ian Wark Research Institute, ARC Special Research Centre, University of
     South Australia, 5095, Australia
SO
     Chemistry of Materials (2005), 17(5), 1027-1035
     CODEN: CMATEX; ISSN: 0897-4756
PΒ
     American Chemical Society
DT
     Journal
LΑ
     English
AB
     We herein report the synthesis of organic-inorg. hybrid poly (Me
     methacrylate) containing polyhedral oligosilsesquioxanes.
     Octakis(3-hydroxypropyldimethylsiloxy)octasilsesquioxane (OHPS) was
     synthesized from octakis(hydridodimethylsiloxy)octasilsesquioxane
     [Si8012 (OSiMe2H) 8, Q8M8H] following literature procedures.
     Octakis (methacryloxypropyldimethylsiloxy) octasilsesquioxane (OMPS) was
     synthesized via the reaction of methacryloyl chloride or methacrylic acid
     anhydride with OHPS, with the latter giving improved purity. Polymerization of
     OMPS with Me methacrylate using a dibenzoyl peroxide initiator gave a
     highly cross-linked polymer. Characterization of the polymer was
    performed using Fourier transform IR spectroscopy, 29Si NMR, differential
     scanning calorimetry, thermogravimetric anal., atomic force microscopy, and
     transmission electron microscopy with energy-dispersive X-ray anal. The
    polymer was found to be largely homogeneous. Increasing the OMPS concentration
in
     the polymer gave increased decomposition and glass transition temps.
CC
     35-6 (Chemistry of Synthetic High Polymers)
ST
     org inorg polymethacrylate hybrid confined polyhedral
     oligosilsesquioxane macromonomer
IT
     Glass transition temperature
       Hybrid organic-inorganic materials
     Polymer morphology
     Thermal stability
        (organic-inorg. poly(Me methacrylate) hybrids with confined
        polyhedral oligosilsesquioxane macromonomers)
ΙT
     849111-73-5P
    RL: PRP (Properties); SPN (Synthetic preparation); PREP
```

(Preparation) (organic-inorg. poly(Me methacrylate) hybrids with confined polyhedral oligosilsesquioxane macromonomers) IT 288290-34-6P RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent) (preparation and polymerization with Me methacrylate) IT 288290-32-4P RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent) (preparation and reaction with methacryloyl chloride or methacrylic acid anhydride) IT 125756-69-6 RL: RCT (Reactant); RACT (Reactant or reagent) (reaction with allyl alc.) IT 107-18-6, Allyl alcohol, reactions RL: RCT (Reactant); RACT (Reactant or reagent) (reaction with octakis(dimethylsiloxy)pentacyclooctasiloxane) IT 760-93-0, Methacrylic acid anhydride 920-46-7, Methacryloyl chloride RL: RCT (Reactant); RACT (Reactant or reagent) (reaction with octakis(hydroxypropyldimethylsilyl)octasilsesquioxane) IT 849111-73-5P RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (organic-inorg. poly(Me methacrylate) hybrids with confined polyhedral oligosilsesquioxane macromonomers) RN849111-73-5 HCAPLUS CN 2-Propenoic acid, 2-methyl-, pentacyclo[9.5.1.13,9.15,15.17,13]octasiloxan e-1,3,5,7,9,11,13,15-octayloctakis[oxy(dimethylsilylene)-3,1-propanediyl] ester, polymer with methyl 2-methyl-2-propenoate (9CI) (CA INDEX NAME) CM 288290-34-6 CRN C72 H136 O36 Si16

$$^{
m CH_2}_{||}$$
 Me $^-$ C $^-$ R

$$\begin{array}{ccc} \text{H}_2\text{C} & \text{O} \\ & || & || \\ \text{Me-C-C-OMe} \end{array}$$

IT 288290-34-6P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent) (preparation and polymerization with Me methacrylate)

RN 288290-34-6 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, pentacyclo[9.5.1.13,9.15,15.17,13]octasiloxan e-1,3,5,7,9,11,13,15-octayloctakis[oxy(dimethylsilylene)-3,1-propanediyl] ester (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 2-B

 $Si-(CH_2)_3-O-C-C-Me$

PAGE 3-B

THERE ARE 23 CITED REFERENCES AVAILABLE FOR THIS RECORD RE.CNT 23 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L37 ANSWER 11 OF 27/ HCAPLUS COPYRIGHT 2007 ACS on STN

AN · 2005:74164 HCAPLUS

DN 142:178756

TI Polyurethane-polymer hybrid dispersion with enhanced surface properties, method for the production and utilization thereof.

Maier, Alois; Ingrisch, Stefan; Steidl, Norbert; Weinelt, Frank IN

Construction Research & Technology G.m.b.H., Germany PA

SO PCT Int. Appl., 46 pp.

CODEN: PIXXD2

DTPatent

LA German

FAN.CNT 1																			
		PATENT NO.					KIN	D :	DATE		APPLICATION NO.						DATE		
I	PΙ	WO 2005007762				A1 20050127			WO 2004-EP7592					20040709					
			W:	ΑE,	AG,	AL,	AM,	AT,	AU,	AZ,	BA,	BB,	BG,	BR,	BW,	BY,	BZ.	CA,	CH,
																			GD,
										IL,									
										MA,									
										PT,									
					-	-	-				-			-	•		•	•	ZW
			RW:	BW,															
							-		-	TJ,	-	•	•	• .	•		•	•	•
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										CG,									
				SNI	TD	TG,	Br,	ъ,	CI,	CG,	CI,	CM,	GA,	GIV,	GΩ,	GW,	ил,	PIK,	ME,
		SN, TD, TG			31 00050303				DE 2002 10221404					00000000					
									DE 2003-10331484										
		EP 1656428			A1 20060517				EP 2004-740871						20040709				
			R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	IT,	LI,	LU,	NL,	SE,	MC,	PT,
										BG,									
		US 2006189750				A1	A1 20060824			US 2006-563903						20060106			
Ε	PRAI	DE 2003-10331484				4	Α												
								W 20040709											
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AΒ Antisoiling, water-resistant, one- and two-component coating materials based on fluorinated polyurethane-polymer hybrid dispersion with enhanced surface properties (low critical surface stresses yc and a

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IT

very high contact angle θ) prepared by (a) producing a dispersion component (binder) based on an aqueous solution or dispersion of an optionally hydroxy and/or amino-functional polyurethane-polymer hybrid with optionally fluorinated side chains and optionally (b) crosslinking the resulting dispersion component. A dispersion of an optionally hydroxy and/or amino-functional polyurethane-polymer is prepared by reacting F-containing anionic-stabilized polyurethane dispersion with (meth)acrylic and aromatic monomers or polymerizing F-containing unsatd. monomers with other (meth)acrylic monomers. Thus, a radical polymerization of a mixture containing 400 g of F-modified polyurethane dispersion (having F-content 0.64 weight% and solid content 38%), 72.12 g of water, 20.27 g of Bu acrylate, 81.07 g of MMA and 1.27 g of an initiator 5 h at 80 - 85° gave a fluorinated polyurethane-polymer hybrid dispersion having solid content 45 weight%. ICM C09D175-04 ICS C08G018-08; C08F283-00 42-7 (Coatings, Inks, and Related Products) antisoiling water resistant coating material fluorinated polyurethane hybrid dispersion; fluorine contg polyurethane dispersion methacrylic monomer polymn dispersion manuf Polyurethanes, uses RL: IMF (Industrial manufacture); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses) (acrylates, fluorine-containing, crosslinked coating; antisoiling, water-resistant coating materials based on fluorinated polyurethane-polymer hybrid dispersion with enhanced surface properties) Coating materials (antisoiling, water-resistant; antisoiling, water-resistant coating materials based on fluorinated polyurethane-polymer hybrid dispersion with enhanced surface properties) Disperse systems (aqueous; antisoiling, water-resistant coating materials based on fluorinated polyurethane-polymer hybrid dispersion with enhanced surface properties) Fluoropolymers, reactions Perfluoro compounds RL: RCT (Reactant); RACT (Reactant or reagent) (fluorinated polyurethane precursor; antisoiling, water-resistant coating materials based on fluorinated polyurethane-polymer hybrid dispersion with enhanced surface properties) Polyurethanes, uses RL: IMF (Industrial manufacture); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses) (perfluoroalkyl group-containing, acrylated; antisoiling, water-resistant coating materials based on fluorinated polyurethane-polymer hybrid dispersion with enhanced surface properties) Fluoropolymers, uses RL: IMF (Industrial manufacture); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses) (polyurethane-, acrylates, crosslinked coating; antisoiling, water-resistant coating materials based on fluorinated polyurethane-polymer hybrid dispersion with enhanced surface properties) Fluoropolymers, uses RL: IMF (Industrial manufacture); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)

(polyurethane-, perfluoroalkyl group-containing, acrylated; antisoiling,

water-resistant coating materials based on fluorinated

polyurethane-polymer **hybrid** dispersion with enhanced surface properties)

IT Acrylic polymers, uses

RL: IMF (Industrial manufacture); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)

(polyurethane-siloxane-, fluorine-containing, crosslinked coating; antisoiling, water-resistant coating materials based on fluorinated polyurethane-polymer hybrid dispersion with enhanced surface properties)

IT 80-62-6DP, Methyl methacrylate, reaction products with perfluoroalkyl group-containing polyurethanes 141-32-2DP, Butyl acrylate, reaction products with perfluoroalkyl group-containing polyurethanes 1996-88-9DP, 3,3,4,4,5,5,6,6,7,7,8,8,9,9,10,10,10-Heptadecafluorodecyl methacrylate, reaction products with perfluoroalkyl group-containing polyurethanes and acrylic monomers 2144-53-8DP, 3,3,4,4,5,5,6,6,7,7,8,8,8Tridecafluorooctyl methacrylate, reaction products with perfluoroalkyl group-containing polyurethanes and acrylic monomers 307531-94-8DP, reaction products with perfluoroalkyl group-containing polyurethanes and acrylic monomers

RL: IMF (Industrial manufacture); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)

(crosslinked coating; antisoiling, water-resistant coating materials based on fluorinated polyurethane-polymer hybrid dispersion with enhanced surface properties)

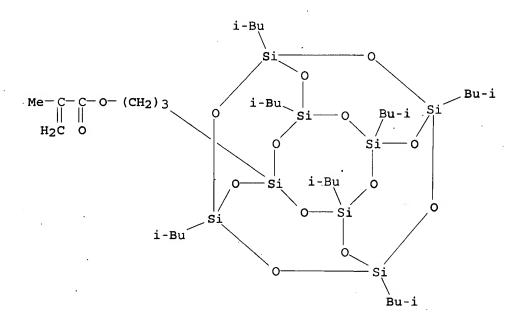
IT 307531-94-8DP, reaction products with perfluoroalkyl group-containing polyurethanes and acrylic monomers

RL: IMF (Industrial manufacture); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)

(crosslinked coating; antisoiling, water-resistant coating materials based on fluorinated polyurethane-polymer hybrid dispersion with enhanced surface properties)

RN 307531-94-8 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-[3,5,7,9,11,13,15-heptakis(2-methylpropyl)pentacyclo[9.5.1.13,9.15,15.17,13]octasiloxan-1-yl]propyl ester (CA INDEX NAME)



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THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
     ANSWER 12 OF 27 HCAPLUS COPYRIGHT 2007 ACS on STN
AN
     2004:874865 HCAPLUS
DN
     142:56733
TI
     Living Radical Polymerization by Polyhedral Oligomeric
     Silsesquioxane-Holding Initiators: Precision Synthesis of Tadpole-Shaped
     Organic/Inorganic Hybrid Polymers
AU
     Ohno, Kohji; Sugiyama, Satoshi; Koh, Kyoungmoo; Tsujii, Yoshinobu; Fukuda,
     Takeshi; Yamahiro, Mikio; Oikawa, Hisao; Yamamoto, Yasuhiro; Ootake,
     Nobumasa; Watanabe, Kenichi
CS
     Institute for Chemical Research, Kyoto University, Uji, Kyoto, 611-0011,
     Japan
SO
     Macromolecules (2004), 37(23), 8517-8522
     CODEN: MAMOBX; ISSN: 0024-9297
PB
     American Chemical Society
     Journal
DT
LΑ
     English
AB
     Incompletely condensed polyhedral oligomeric silsesquioxane (POSS) with
     the highly reactive group of trisodium silanolate was used for the
     synthesis of two initiators for atom transfer radical polymerization, one with
а
     2-bromoisobutyl group and the other with a chlorosulfonyl group.
     initiators were applied to solution polymns. of styrene and Me methacrylate
     in the presence of a copper complex. In both systems, polymerization proceeded
     in a living fashion, as indicated by the first-order kinetics of monomer
     consumption, the evolution of mol. weight in direct proportion to monomer
     conversion, the good agreement of mol. weight with the theor. one, and the
     low polydispersity, thus providing tadpole-shaped polymers with an "inorg.
     head" of POSS and an "organic tail" of well-defined polymer.
     Thermogravimetric and differential scanning calorimetric studies showed
     that both thermal degradation and glass transition temps. of the organic/inorg.
     hybrid polymers with mol. wts. up to about 20 000 were enhanced as
     compared to those of model polymers without the POSS moiety.
     35-4 (Chemistry of Synthetic High Polymers)
     polyhedral oligomeric silsesquioxane initiator prepn ATRP polymn
st
IT
     Polymerization catalysts
        (atom transfer, living, radical; living radical polymerization by polyhedral
        oligomeric silsesquioxane-holding initiators: precision synthesis of
        Tadpole-shaped organic/inorg. hybrid polymers)
     Hybrid organic-inorganic materials
        (living radical polymerization by polyhedral oligomeric
silsesquioxane-holding
        initiators: precision synthesis of Tadpole-shaped organic/inorg.
        hybrid polymers)
IT
     Silsesquioxanes
     RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation);
     USES (Uses)
        (living radical polymerization by polyhedral oligomeric
silsesquioxane-holding
        initiators: precision synthesis of Tadpole-shaped organic/inorg.
        hybrid polymers)
TΤ
     Glass transition temperature
     Molecular weight
     Polydispersity
        (of Tadpole-shaped organic/inorg. hybrid polymers)
IT
     Polymer degradation
        (temperature; of Tadpole-shaped organic/inorg. hybrid polymers)
```

660392-78-9P

IT

PENG 10/520763 07/11/2007Page 104

RL: CAT (Catalyst use); RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)

(in preparation of living radical polymerization polyhedral oligomeric silsesquioxane-holding/initiators)

IT 18204-80-3 20769-85-1 79793-00-3 476635-00-4

RL: RCT (Reactant); RACT (Reactant or reagent)

(in preparation of living radical polymerization polyhedral oligomeric silsesquioxane-holding initiators)

IT 757198-90-6P 757199-15-8P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(in preparation of living radical polymerization polyhedral oligomeric silsesquioxane-holding initiators)

IT 757199-48-7P

RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP
(Preparation); USES (Uses)

(preparation of living radical polymerization polyhedral oligomeric silsesquioxane-holding initiators)

IT 9003-53-6P, Polystyrene 9011-14-7P, Poly(methyl methacrylate)

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (synthesis of Tadpole-shaped organic/inorg. hybrid polymers

catalyzed by polyhedral oligomeric silsesquioxane-holding initiators)

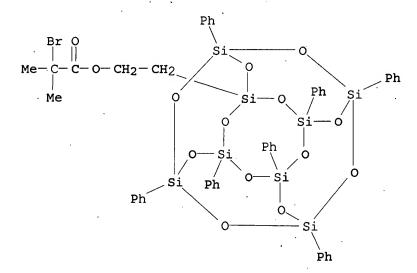
IT. 757199-48-7P

RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)

(preparation of living radical polymerization polyhedral oligomeric silsesquioxane-holding initiators)

RN 757199-48-7 HCAPLUS

CN Propanoic acid, 2-bromo-2-methyl-, 2-(3,5,7,9,11,13,15heptaphenylpentacyclo[9.5.1.13,9.15,15.17,13]octasiloxanyl)ethyl ester (9CI) (CA INDEX NAME)



RE.CNT 41 THERE ARE 41 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L37 ANSWER 13 OF 27 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2004:230854 HCAPLUS

DN 141:54760

TI Ultrathin films of self-assembled organic-inorganic hybrid

nanoparticle block copolymers

AU Intasanta, Narupol; Russell, Thomas P.; Coughlin, E. Bryan

CS Department of Polymer Science and Engineering, University of

Massachusetts, Amherst, MA, 01003, USA

SO PMSE Preprints (2004), 90, 260-261

CODEN: PPMRA9; ISSN: 1550-6703

PB American Chemical Society

DT Journal; (computer optical disk)
LA English

AB [3-(3,5,7,9,11,13,15-Heptaisobutylpentacyclo-9.5.1.13,9.15,15.17,13)octasi loxan-1-yl]propyl methacrylate [poly(methacrylate-polyhedral oligomeric silsesquioxane (POSS)-(isobutyl)-b-Me methacrylate)] hybrids of mass fraction 60:40 based on PMMA:POSS(i-Butyl), with a PDI of 1.3, and number average mol. weight of 40k was prepared by atom transfer radical polymerization The

preparation of P[MAPoss(1-butyl)-b-MMA] was possible by the sequential polymerization

of the inorg. block first followed by the organic domain. Small angle x-ray scattering (SAXS) data show that these hybrid diblock copolymers have a long period spacing of 25 nm. Studies by atomic force microscopy clearly show cylindrical morphol. The cylindrical domains lie parallel to the substrate at the air/polymer interface. Modification of the silicon oxide sub-surface with a neutral brush copolymer results in hybrid block copolymer cylinders oriented perpendicular to the substrate. UV irradiation results in selective decomposition of the PMMA domain and allows

for removal of the parallel layer by acetic acid rinse. The cylinders oriented perpendicular to the surface undergo oxygen plasma conversion of the silsesquioxane domains into nano-patterned silicon oxide nanoarrays.

CC 35-8 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 36, 76

ST methyl methacrylate cyclooctasiloxane propyl hybrid block copolymer prepn; self assembly hybrid cyclosiloxane methacrylate block copolymer morphol orientation; silsesquioxane domain oxygen plasma reaction silica array nanopattern array

IT Polymer morphology

(phase; preparation of silica nanoarrays by decomposition of prepared ultrathin

films of self-assembled **hybrid** cyclosiloxane-methacrylate block copolymer nanoparticles)

IT Hybrid organic-inorganic materials

Molecular orientation

Self-assembly

(preparation of silica nanoarrays by decomposition of prepared ultrathin films of

self-assembled hybrid cyclosiloxane-methacrylate block
copolymer nanoparticles)

IT Polymer degradation

(radiochem.; preparation of silica nanoarrays by decomposition of prepared ultrathin films of self-assembled hybrid cyclosiloxane-methacrylate block copolymer nanoparticles)

IT Nanoparticles

(silica nanoarrays; preparation of silica nanoarrays by decomposition of prepared

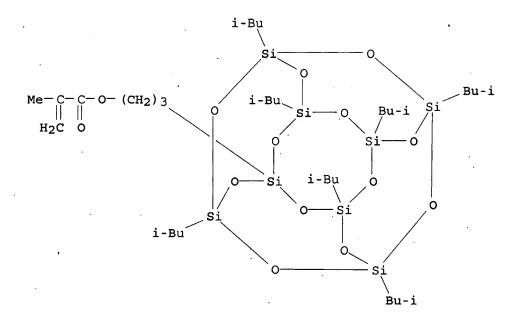
ultrathin films of self-assembled hybrid cyclosiloxanemethacrylate block copolymer nanoparticles)

IT 25034-86-0, Methyl methacrylate-styrene copolymer RL: NUU (Other use, unclassified); USES (Uses)

(brush polymer, surface layer on substrate; preparation of silica nanoarrays by decomposition of prepared ultrathin films of self-assembled hybrid

CRN 307531-94-8 C35 H74 O14 Si8 CMF

PENG



CRN 80-62-6 CMF C5 H8 O2

RE.CNT 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L37 ANSWER 14 OF 27 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2003:384332 HCAPLUS

DN 139:180690

TI Organic-inorganic hybrid gels having functionalized silsesquioxanes

AU Kim, Kyung-Min; Chujo, Yoshiki

CS Department of Polymer Chemistry, Graduate School of Engineering, Kyoto University, Kyoto, 606-8501, Japan

SO Journal of Materials Chemistry (2003), 13(6), 1384-1391 CODEN: JMACEP; ISSN: 0959-9428

PB Royal Society of Chemistry

DT Journal

LA English

AB Hybrid terpolymers consisting of polyhedral oligomeric silsesquioxanes (POSS), N,N-dimethylacrylamide, and bipyridine monomer were synthesized by a common radical polymerization method and their structures and thermal properties were studied by FT-IR, 1H NMR, DSC, TGA. The thermal stabilities of the hybrid terpolymers increased on increasing the content of POSS in the feed ratio. New hybrid gels containing POSS were prepared through the coordination of various metal ions to 2,2'-bipyridine-modified hybrid terpolymers. Highly

CC

ST

IT

IT

IT

IT

IT.

IT

TΤ

TT

IT

concentrated solns. of terpolymers with iron(ii) sulfate or ruthenium(iii) chloride gave hybrid gels in good yields. No hybrid gel was formed with nickel(ii) chloride even at much higher concns. of nickel ions due to rapid ligand exchange reactions causing the polymer network to disappear as a result of a change in the nature of the coordination bonds from an intermol. network to an intramol. network. degree of swelling and the thermal stability of the gels in various solvents were dependent on the content of POSS moiety in the hybrid gel and the ruthenium gel was considerably more stable than the iron and nickel gels. The hybrid gels containing POSS had properties characteristic of hydrogels and those of lipogel depending on the content of POSS in the hybrid gel. By anal. of the degree of swelling, the amount of POSS moiety in the hybrid gel has a significant effect on the degree of swelling in this system. 37-3 (Plastics Manufacture and Processing) functionalized silsesquioxane terpolymer prepn org inorg hybrid Silsesquioxanes RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PRP (Properties); SPN (Synthetic preparation); PREP (Preparation); PROC (Process) (acrylic; preparation of organic-inorg. hybrid gels from terpolymer containing functionalized silsesquioxanes in the presence of metal ions) Differential thermal analysis Polymer morphology Swelling, physical Thermal stability Thermogravimetric analysis (of hybrid gels formed from terpolymer containing functionalized silsesquioxanes in the presence of metal ions) Glass transition temperature (of organic-inorg. hybrid terpolymer containing functionalized silsesquioxanes and hybrid gels therefrom in the presence of metal ion) Gelation (of organic-inorg. hybrid terpolymer containing functionalized silsesquioxanes in the presence of metal ion) Hybrid organic-inorganic materials (preparation of organic-inorg. hybrid gels from terpolymer containing functionalized silsesquioxanes in the presence of metal ions) 95314-35-5P 183387-28-2P RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent) (in preparation of hybrid terpolymer containing functionalized silsesquioxanes for organic-inorg. gels) 74173-48-1P 387820-81-7P RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent) (monomer; in preparation of hybrid terpolymer containing functionalized silsesquioxanes for organic-inorq. qels) 579470-10-3P RL: PRP (Properties); SPN (Synthetic preparation); PREP (preparation of hybrid terpolymer containing functionalized silsesquioxanes for organic-inorq. hybrid gels) 7439-89-6DP, Iron, complex with silsesquioxane- and bipyridine-containing 7440-02-0DP, Nickel, complex with silsesquioxane- and bipyridine-containing terpolymer 7440-18-8DP, Ruthenium, complex with silsesquioxane- and bipyridine-containing terpolymer 579470-10-3DP,

complex with metal ions

RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PRP (Properties); SPN (Synthetic preparation); PREP (Preparation); PROC (Process)

(preparation of organic-inorg. hybrid gels from terpolymer containing functionalized silsesquioxanes in the presence of metal ions)

IT

1134-35-6, 4,4'-Dimethyl-2,2'-bipyridine 14579-03-4, Cyclopentyltrichlorosilane 38595-89-0, (3-Acryloxypropyl)trichlorosilane RL: RCT (Reactant); RACT (Reactant or reagent)
(reactant for monomer; in preparation of hybrid terpolymer containing

functionalized silsesquioxanes for organic-inorg. gels)

IT 387820-81-7P

> RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(monomer; in preparation of hybrid terpolymer containing functionalized silsesquioxanes for organic-inorg. gels)

RN387820-81-7 HCAPLUS

CN 2-Propenoic acid, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,15.17,13]oct asiloxanyl)propyl ester (9CI) (CA INDEX NAME)

IT 579470-10-3P

RL: PRP (Properties); SPN (Synthetic preparation); PREP

(preparation of hybrid terpolymer containing functionalized silsesquioxanes for organic-inorg. hybrid gels)

RN 579470-10-3 HCAPLUS

> 2-Propenoic acid, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,15.17,13]oct asiloxanyl)propyl ester, polymer with N,N-dimethyl-2-propenamide and 4-ethenyl-4'-methyl-2,2'-bipyridine (9CI) (CA INDEX NAME)

CM

CN

387820-81-7 CMF C41 H72 O14 Si8

CRN 74173-48-1 CMF C13 H12 N2

CM 3

CRN 2680-03-7 CMF C5 H9 N O

$$\begin{array}{c} & \text{O} \\ || \\ \text{Me}_2 \text{N-C-CH-} \end{array} \text{CH}_2$$

PENG 10/520763 07/11/2007Page 111

RN 579470-10-3 HCAPLUS

CN 2-Propenoic acid, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,15.17,13]oct asiloxanyl)propyl ester, polymer with N,N-dimethyl-2-propenamide and 4-ethenyl-4'-methyl-2,2'-bipyridine (9CI) (CA INDEX NAME)

CM 1

CRN 387820-81-7 CMF C41 H72 O14 Si8

CM 2

CRN 74173-48-1 CMF C13 H12 N2

CM 3

CRN 2680-03-7 CMF C5 H9 N O

```
Me_2N-C-CH=CH_2
              THERE ARE 37 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT 37
              ALL CLATIONS AVAILABLE IN THE RE FORMAT
     ANSWER 15 OF 27 HCAPLUS COPYRIGHT 2007 ACS on STN
L37
AN
     2002:716627 HCAPLUS
DN
     137:270509
    High resolution resists comprising nanoparticles and inorganic moieties
TI
     for next generation lithographies
IN
     Gonsalves, Kenneth E.
PA
     University of North Carolina at Charlotte, USA; University of Connecticut
     PCT Int. Appl., 62 pp.
so
     CODEN: PIXXD2
DT
     Patent
LA
    English
FAN.CNT 4
     PATENT NO.
                         KIND
                                DATE
                                            APPLICATION NO.
                         _ _ _ _
                                           -----
PΙ
    WO 2002073308
                         A1
                                20020919 WO 2002-US7338
                                                                   20020311
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
             PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ,
             UA, UG, US, UZ, VN, YU, ZA, ZM, ZW-
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH,
             CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR,
             BF, BJ, CF, CG, CI, CM, GA, GN, GO, GW, ML, MR, NE, SN, TD, TG
    AU 2002254170
                          A1
                                20020924
                                          AU 2002-254170
                                                                   20020311
     EP 1377876
                         A1
                                20040107
                                           EP 2002-723388
                                                                   20020311
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
```

IE, SI, LT, LV, FI, RO, MK, CY, AL, TR

20041007

20010312

20020311

JP 2002-572502

20020311

Т

Р

W

JP 2004530921

WO 2002-US7338

PRAI US 2001-274719P

GI

AB The present invention provides new high resolution resists applicable to next generation lithogs., methods of making these novel resists, and methods of using these new resists in lithog. processes to effect state-of-the-art lithogs. New nanocomposite resists comprising polymers of the general formula I (R = alkyl, cycloalkyl, silyl, aryl, aralkyl, alkenyl) and nanoparticles in a polymer matrix are provided in the invention. New chemical amplified resists that incorporate inorg. moieties as part of the polymer and chemical amplified resists that incorporate photoacid generating groups within the polymeric chain are presented. Novel non-chemical amplified yet photosensitive resists, and new organic-inorg. hybrid resists are also provided. This invention and the embodiments described constitute fundamentally new architectures for high resolution resists that achieve high sensitivity, contrast, resolution and high plasma etch resistance.

I

IC ICM G03C001-725

ICS G03F007-039; G03F007-075; G03F007-26

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
Section cross-reference(s): 38, 76

ST chem amplified resist nanoparticle silsesquioxane photoacid generator copolymer polymer; lithog electron ion beam x ray chem amplified resist; photolithog UV chem amplified resist nanoparticle silsesquioxane

IT Photolithography

(UV; chemical amplified resists comprising copolymers with sulfonium photoacid generator monomer for)

IT Resists

(chemical amplified resists comprising copolymers with sulfonium photoacid generator monomer)

IT Electron beam lithography

Ion beam lithography

X-ray lithography

(chemical amplified resists comprising copolymers with sulfonium photoacid generator monomer for)

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IT
      Integrated circuits
         (chemical amplified resists comprising copolymers with sulfonium photoacid
         generator monomer for fabrication of)
IT
      Polyoxymethylenes, preparation
      RL: PNU (Preparation, unclassified); TEM (Technical or engineered material
     use); PREP (Preparation); USES (Uses)
         (chemical amplified resists comprising polyacetals)
IT
      Silsesquioxanes
      RL: TEM (Technical or engineered material use); USES (Uses)
         (chemical amplified resists comprising polyhydral oligosilsesquioxanes,
         nanoparticles and inorg. moieties)
      43127-35-1, ZEP 520
IT
      RL: TEM (Technical or engineered material use); USES (Uses)
         (ZEP 520; chemical amplified resists comprising polyhydral
         oligosilsesquioxanes, nanoparticles and inorg. moieties)
IT
      352455-55-1P 362675-17-0P 461699-74-1P
      RL: PRP (Properties); SPN (Synthetic preparation); TEM (Technical or
      engineered material use); PREP (Preparation); USES (Uses)
         (chemical amplified resists comprising copolymers with sulfonium photoacid
         generator monomer)
IT
      461699-77-4P
                     461699-80-9P
     RL: PNU (Preparation, unclassified); TEM (Technical or engineered material
     use); PREP (Preparation); USES (Uses)
         (chemical amplified resists comprising polyacetals)
IT
     359408-40-5P
     RL: PRP (Properties); SPN (Synthetic preparation); TEM (Technical or
      engineered material use); PREP (Preparation); USES (Uses)
         (chemical amplified resists comprising polyhydral oligosilsesquioxanes,
         nanoparticles and inorg. moieties)
ΙT
     136849-03-1
     RL: TEM (Technical or engineered material use); USES (Uses)
         (chemical amplified resists comprising polyhydral oligosilsesquioxanes,
         nanoparticles and inorg. moieties)
     338731-99-0P
     RL: PRP (Properties); SPN (Synthetic preparation); TEM (Technical or
      engineered material use); PREP (Preparation); USES (Uses)
         (chemical amplified resists comprising sulfonium photoacid generator
IT
     2170-03-8, Itaconic anhydride
     RL: TEM (Technical or engineered material use); USES (Uses)
         (dissoln. promoter; chemical amplified resists comprising copolymers with
         sulfonium photoacid generator monomer)
IT
     352455-54-0P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
      (Reactant or reagent)
         (in preparation of copolymers containing sulfonium photoacid generator
monomer)
                                    920-46-7, Methacryloyl chloride
IT
     108-95-2, Phenol, reactions
     RL: RCT (Reactant); RACT (Reactant or reagent)
         (in preparation of sulfonium photoacid generator monomer)
IT
                   301152-82-9P
                                  364325-13-3P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
      (Reactant or reagent)
         (in preparation of sulfonium photoacid generator monomer)
IT
     352455-55-1P 362675-17-0P 461699-74-1P
     RL: PRP (Properties); SPN (Synthetic preparation); TEM (Technical or
     engineered material use); PREP (Preparation); USES (Uses)
         (chemical amplified resists comprising copolymers with sulfonium photoacid
        generator monomer)
     352455-55-1 HCAPLUS
```

CN Sulfonium, dimethyl[4-[(2-methyl-1-oxo-2-propenyl)oxy]phenyl]-, salt with trifluoromethanesulfonic acid (1:1), polymer with 1,1-dimethylethyl 2-methyl-2-propenoate and 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,15.1 7,13]octasiloxanyl)propyl 2-methyl-2-propenoate (9CI) (CA INDEX NAME)

CM 1

CRN 169391-91-7 CMF C42 H74 O14 Si8

CM 2

CRN 585-07-9 CMF C8 H14 O2

$$\begin{array}{c|c} \text{O} & \text{CH}_2 \\ \parallel & \parallel \\ \text{t-BuO-C-C-Me} \end{array}$$

CM 3

CRN 352455-54-0 CMF C12 H15 O2 S . C F3 O3 S

CM 4

CRN 141718-72-1 CMF C12 H15 O2 S

$$\begin{array}{c|c} O & CH_2 \\ \parallel & \parallel \\ O-C-C-Me \\ \\ Me & \\ \end{array}$$

CRN 37181-39-8 CMF C F3 O3 S

RN 362675-17-0 HCAPLUS

CN Sulfonium, dimethyl[4-[(2-methyl-1-oxo-2-propenyl)oxy]phenyl]-, salt with trifluoromethanesulfonic acid (1:1), polymer with dihydro-3-methylene-2,5-furandione, 1,1-dimethylethyl 2-methyl-2-propenoate, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,15.17,13]octasiloxanyl)propyl 2-methyl-2-propenoate and methyl 2-methyl-2-propenoate (9CI) (CA INDEX NAME)

CM 1

CRN 169391-91-7 CMF C42 H74 O14 Si8

CRN 2170-03-8 CMF C5 H4 O3

CM 3

CRN 585-07-9 CMF C8 H14 O2

$$\begin{array}{c|c} \text{O} & \text{CH}_2 \\ \parallel & \parallel \\ \text{t-BuO-C-C-Me} \end{array}$$

CM 4

CRN 80-62-6 CMF C5 H8 O2

$$\begin{array}{ccc} ^{H_2C} & \text{O} \\ & \parallel & \parallel \\ \text{Me-} & \text{C-} & \text{C-} & \text{OMe} \end{array}$$

CRN 352455-54-0 CMF C12 H15 O2 S . C F3 O3 S

CM 6

CRN 141718-72-1 CMF C12 H15 O2 S .

CM 7

CRN 37181-39-8 CMF C F3 O3 S

CN

RN 461699-74-1 HCAPLUS

Sulfonium, dimethyl[4-[(2-methyl-1-oxo-2-propenyl)oxy]phenyl]-, salt with trifluoromethanesulfonic acid (1:1), polymer with 1,1-dimethylethyl 2-methyl-2-propenoate, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,15.17,1 3]octasiloxanyl)propyl 2-methyl-2-propenoate, methyl 2-methyl-2-propenoate and 2-methyl-2-propenoic acid (9CI) (CA INDEX NAME)

CM 1

CRN 169391-91-7 CMF C42 H74 O14 Si8

CRN 585-07-9 CMF C8 H14 O2

$$\begin{array}{c|c} & \text{O} & \text{CH}_2 \\ & || & || \\ \text{t-BuO-C-C-Me} \end{array}$$

CM 3

CRN 80-62-6 CMF C5 H8 O2

$$\begin{array}{ccc} ^{\text{H}_2\text{C}} & \text{O} \\ \parallel & \parallel \\ ^{\text{Me-}} \text{C--C-OMe} \end{array}$$

CM 4

CRN 79-41-4 CMF C4 H6 O2

$$\begin{array}{c} \text{CH}_2 \\ || & \cdot \\ \text{Me-} \text{C-} \text{CO}_2 \text{H} \end{array}$$

CRN 352455-54-0 CMF C12 H15 O2 S . C F3 O3 S

CM 6

CRN 141718-72-1 CMF C12 H15 O2 S

CM 7

CRN 37181-39-8 CMF C F3 O3 S

CN

IT 359408-40-5P

RL: PRP (Properties); SPN (Synthetic preparation); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses) (chemical amplified resists comprising polyhydral oligosilsesquioxanes, nanoparticles and inorq. moieties)

RN 359408-40-5 HCAPLUS

2-Propenoic acid, 2-methyl-, polymer with 1,1-dimethylethyl 2-methyl-2-propenoate, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,15.17,1 3]octasiloxanyl)propyl 2-methyl-2-propenoate and methyl 2-methyl-2-propenoate (9CI) (CA INDEX NAME)

CM 1

CRN 169391-91-7 CMF C42 H74 O14 Si8 PENG 10/520763 07/11/2007Page 121

CM 2

CRN 585-07-9 CMF C8 H14 O2

$$\begin{array}{c|c} \text{O} & \text{CH}_2 \\ \parallel & \parallel \\ \text{t-BuO-C-C-Me} \end{array}$$

CM 3

CRN 80-62-6 CMF C5 H8 O2

$$\begin{array}{c|c} ^{H_2C} & \text{O} \\ \parallel & \parallel \\ \text{Me-} \text{C-} \text{C-} \text{OMe} \end{array}$$

CM 4

CRN 79-41-4 CMF C4 H6 O2

$$\begin{array}{c} \text{CH}_2 \\ || \\ \text{Me-} \text{C-} \text{CO}_2 \text{H} \end{array}$$

PENG 10/520763 07/11/2007Page 122 THERE ARE A CITED REFERENCES AVAILABLE FOR THIS RECORD RE.CNT ALL CITATIONS AVAILABLE IN THE RE FORMAT ANSWER 16 OF 27 HCAPLUS COPYRIGHT 2007 ACS on STN L37 2002:483838 HCAPLUS AN DN 137:248272 TI Synthesis and Properties of Perfluoroalkyl Groups Containing Double Four-Ring Spherosilicate (Siloxysilsesquioxane) Precursors Hoebbel, Dagobert; Weber, Christine; Schmidt, Helmut; Krueger, Ralph-Peter ΑU Institut fuer Neue Materialien GmbH, Saarbruecken, D-66123, Germany CS SO Journal of Sol-Gel Science and Technology (2002), 24(2), 121-129 CODEN: JSGTEC; ISSN: 0928-0707 PB Kluwer Academic Publishers DT Journal LA English AB Organically modified cage-like double four-ring spherosilicates have received considerable interest in the construction of nanosized hybrid materials, as well as building units for structural well-defined polymers. This group is extended by perfluoroalkyl ligands containing spherosilicates, synthesized by addition reaction of the octahydridodimethylsiloxyoctasilsesquioxane [H(CH3)2Si]8Si8O20 and heptadecafluorodecyl methacrylate. The resultant liquid spherosilicate substituted with eight terminal perfluoroalkyl groups was characterized by 29Si and 13C NMR spectroscopies and MALDI Time-of-Flight mass spectrometry. Partial substitution of perfluoroalkyl ligands by trimethoxysilyl containing groups provides condensable precursors for the synthesis of hydrophobic and oleophobic materials via the sol-gel process. This new spherosilicate, carrying on average four perfluoroalkyl groups and four trimethoxysilyl groups shows better hydrophobic and oleophobic properties compared with commonly used perfluoroalkyltrialkoxysilanes under identical concentration of perfluoroalkyl chains. In addition a comprehensive literature survey is given on structural well characterized, organically modified cage-like double four-ring spherosilicates. 37-3 (Plastics Manufacture and Processing) CC Section cross-reference(s): 57 fluorine contg silsesquioxane synthesis sol ceramic coating hydrophobicity ST TT Surface tension (of coatings; synthesis and characterization of perfluoroalkyl group-containing siloxysilsesquioxanes and their sol coatings) IT Sol-gel processing (polymerization; synthesis and characterization of perfluoroalkyl group-containing siloxysilsesquioxanes and their sol coatings) TT Silsesquioxanes RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (silicate-, fluorine-containing; synthesis and characterization of perfluoroalkyl group-containing siloxysilsesquioxanes and their sol coatings) ΙT Fluoropolymers, preparation RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (silicate-silsesquioxane-; synthesis and characterization of perfluoroalkyl group-containing siloxysilsesquioxanes and their sol

IT Polymerization

(sol-gel; synthesis and characterization of perfluoroalkyl group-containing siloxysilsesquioxanes and their sol coatings)

IT Ceramers

Ceramic coatings

coatings)

(synthesis and characterization of perfluoroalkyl group-containing

```
siloxysilsesquioxanes and their sol coatings)
TΤ
     1996-88-9 2530-85-0, 3-(Methacryloxy) propyltrimethoxysilane
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (reaction with octahydrosilisesquioxane; synthesis and characterization
        of perfluoroalkyl group-containing siloxysilsesquioxanes and their sol.
        coatings)
     125756-69-6
IT
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (reaction with perfluoroalkyl methacrylates; synthesis and
        characterization of perfluoroalkyl group-containing siloxysilsesquioxanes
        and their sol coatings)
IT
     460730-46-5P
                    460991-50-8P
     RL: PRP (Properties); SPN (Synthetic preparation); PREP
     (Preparation)
        (synthesis and characterization of perfluoroalkyl group-containing
        siloxysilsesquioxanes and their sol coatings)
IT
     460730-47-6P 460730-48-7P
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (synthesis and characterization of perfluoroalkyl group-containing
        siloxysilsesquioxanes and their sol coatings)
TΤ
     460730-46-5P
     RL: PRP (Properties); SPN (Synthetic preparation); PREP
     (Preparation)
        (synthesis and characterization of perfluoroalkyl group-containing
        siloxysilsesquioxanes and their sol coatings)
RN
     460730-46-5 HCAPLUS
     Propanoic acid, 3,3',3'',3''',3'''',3'''',3''''',3'''''-
CN
     [pentacyclo[9.5.1.13,9.15,15.17,13]octasiloxane-1,3,5,7,9,11,13,15-
     octayloctakis[oxy(dimethylsilylene)]]octakis[2-methyl-, tetraester with
     3,3,4,4,5,5,6,6,7,7,8,8,9,9,10,10,10-heptadecafluoro-1-decanol tetraester
     with 3-(trimethoxysilyl)-1-propanol (9CI) (CA INDEX NAME)
     CM
     CRN
         460730-45-4
    CMF C48 H104 O36 Si16
```

PAGE 1-A

PAGE 1-B

— со2н

CM 2

CRN 53764-54-8 CMF C6 H16 O4 Si

$$\begin{array}{c} \text{OMe} \\ \cdot & | \\ \text{MeO-Si-} (\text{CH}_2)_3 - \text{OH} \\ | \\ \text{OMe} \end{array}$$

CM 3

CRN 678-39-7 CMF C10 H5 F17 O

```
HO-CH_2-CH_2-(CF_2)_7-CF_3
IT
     460730-47-6P 460730-48-7P
     RL: SPN (Synthetic preparation); PREP (Preparation) (synthesis and characterization of perfluoroalkyl group-containing
        siloxysilsesquioxanes and their sol coatings)
RN
     460730-47-6 HCAPLUS
CN
     [pentacyclo[9.5.1.13,9.15,15.17,13]octasiloxane-1,3,5,7,9,11,13,15-
     octayloctakis[oxy(dimethylsilylene)]]octakis[2-methyl-,
     tetrakis(3,3,4,4,5,5,6,6,7,7,8,8,9,9,10,10,10-heptadecafluorodecyl)
     tetrakis[3-(trimethoxysilyl)propyl] ester, polymer with silicic acid
     (H4SiO4) tetraethyl ester and triethoxymethylsilane (9CI)
                                                               (CA INDEX NAME)
     CM
     CRN
         2031-67-6
     CMF
          C7 H18 O3 Si
     OEt
EtO-Si-Me
     OEt
     CM
          2
     CRN
          78-10-4
     CMF
          C8 H20 Q4 Si
     OEt
Eto-Si-OEt
     OEt
     CM
          3
     CRN 460730-46-5
         C112 H172 F68 O48 Si20
     CMF
     CCI
          IDS
```

460730-45-4

C48 H104 O36 Si16

CM

CRN

CMF

PAGE 1-B

— со2н

CM ·5

CRN 53764-54-8 CMF C6 H16 O4 Si

$$\begin{array}{c} \text{OMe} \\ | \\ \text{MeO-Si-(CH}_2)_3 - \text{OH} \\ | \\ \text{OMe} \end{array}$$

CM 6

CRN 678-39-7 CMF C10 H5 F17 O ${\rm HO-CH_2-CH_2-(CF_2)_{7}-CF_3}$

CM 1

CRN 460730-46-5

CMF C112 H172 F68 O48 Si20

CCI IDS

CM 2

CRN 460730-45-4

CMF C48 H104 O36 Si16

— co2н

CM 3

CRN 53764-54-8 CMF C6 H16 O4 Si

$$\begin{array}{c} \text{OMe} \\ | \\ \text{MeO-Si- (CH2)}_3 - \text{OH} \\ | \\ \text{OMe} \end{array}$$

CM 4

CRN 678-39-7 CMF C10 H5 F17 O

 $HO-CH_2-CH_2-(CF_2)_7-CF_3$

RE.CNT 49 THERE ARE 49 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L37 ANSWER 17 OF 27 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2002:232280 HCAPLUS

DN 136:370435

TI Thermal properties and morphology of POSS/PMMA hybrids prepared by copolymerization and blending

AU Xiao, Jun; Feher, Frank J.

CS Dep. Chem., Univ. California, Irvine, CA, 92697-2025, USA

SO PMSE Preprints (2002), 86, 171-172 CODEN: PPMRA9; ISSN: 1550-6703

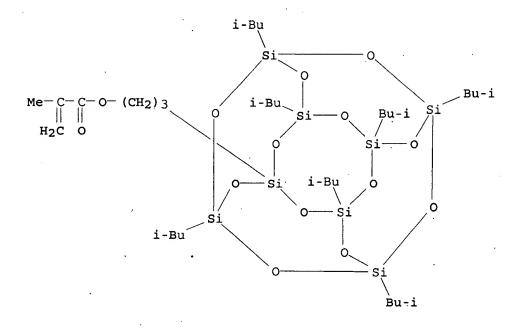
PB · American Chemical Society

DT Journal; (computer optical disk)

LA English

AB Hybrids containing polyhedral oligosilsesquioxanes (POSS) have attracted considerable attention because the incorporation of POSS into traditional organic polymer resins often leads to useful property enhancements. Two common methods used to produce POSS-containing hybrids were copolymn. of functionalized POSS monomers with MMA, and blending of POSS with molten thermoplastics. In this paper, we use both methods to synthesize and characterize POSS/PMMA hybrids containing up to 30 weight % POSS. A detailed anal. of the differential scanning

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calorimetry (DSC) and SEM data indicate that compatibility and miscibility
     play an important role in determining the phys. characteristics of
POSS-containing
     hybrids.
     37-6 (Plastics Manufacture and Processing)
CC
     Section cross-reference(s): 29
     thermal property morphol polyhedral oligosilsesquioxane PMMA
ST
     hybrid
IT
     Polymer morphology
        (fracture-surface; thermal properties and morphol. of POSS/PMMA
        hybrids prepared by copolymn. and blending)
     Silsesquioxanes
IT
     RL: PRP (Properties)
        (oligosilsesquioxanes; thermal properties and morphol. of POSS/PMMA
        hybrids prepared by copolymn. and blending)
IT
     Fracture surface morphology
        (polymeric; thermal properties and morphol. of POSS/PMMA
        hybrids prepared by copolymn. and blending)
     Fusion enthalpy
IT
     Glass transition temperature
       Hybrid organic-inorganic materials
     Miscibility
        (thermal properties and morphol. of POSS/PMMA hybrids prepared
        by copolymn. and blending)
IT
     221326-46-1
     RL: PRP (Properties)
        (mixture with PMMA; thermal properties and morphol. of POSS/PMMA
        hybrids prepared by copolymn. and blending)
IT
     9011-14-7, PMMA
     RL: POF (Polymer in formulation); PRP (Properties); USES (Uses)
        (mixture with oligosilsesquioxane; thermal properties and morphol. of
        POSS/PMMA hybrids prepared by copolymn. and blending)
IT
     425409-07-0P 425409-08-1P
     RL: PRP (Properties); SPN (Synthetic preparation); PREP
     (Preparation)
        (thermal properties and morphol. of POSS/PMMA hybrids prepared
        by copolymn. and blending)
IT
     425409-07-0P 425409-08-1P
     RL: PRP (Properties); SPN (Synthetic preparation); PREP
     (Preparation)
        (thermal properties and morphol. of POSS/PMMA hybrids prepared
        by copolymn. and blending)
RN
     425409-07-0 HCAPLUS
     2-Propenoic acid, 2-methyl-, 3-[3,5,7,9,11,13,15-heptakis(2-
CN
     methylpropyl)pentacyclo[9.5.1.13,9.15,15.17,13]octasiloxan-1-yl]propyl
     ester, polymer with methyl 2-methyl-2-propenoate (CA INDEX NAME)
     CM
         307531-94-8
     CRN
     CMF C35 H74 O14 Si8
```



CRN 80-62-6 CMF C5 H8 O2

$$\begin{array}{c|c} {\rm H_2C} & {\rm O} \\ & || & || \\ {\rm Me-} & {\rm C-} & {\rm C-} & {\rm OMe} \end{array}$$

RN 425409-08-1 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-[heptakis(2-methylpropyl)pentacyclo[9.5.1.1 3,9.15,15.17,13]octasiloxanyl]propyl ester, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 307531-94-8 CMF C35 H74 O14 Si8

RE.CNT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L37 ANSWER 18 OF 27 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2001:806199 HCAPLUS

DN 136:86132

TI Liquid-crystalline organic-inorganic hybrid polymers with functionalized silsesquioxanes

AU Kim, Kyung-Min; Chujo, Yoshiki

CS Department of Polymer Chemistry, Graduate School of Engineering, Kyoto University, Kyoto, 606-8501, Japan

SO Journal of Polymer Science, Part A: Polymer Chemistry (2001), 39(22), 4035-4043
CODEN: JPACEC; ISSN: 0887-624X

John Wiley & Sons, Inc.

DT Journal

PB

LA English

AB Liquid-crystalline (LC) hybrid polymers with functionalized silsesquioxanes with various proportions of LC monomer were synthesized by the reaction of polyhedral oligomeric silsesquioxane (POSS) macromonomer with methacrylate monomer having an LC moiety under common free-radical conditions. The obtained LC hybrid polymers were soluble in common solvents such as THF, toluene, and chloroform, and their structures were characterized with Fourier transform IR, 1H NMR, and 29Si NMR. thermal stability of the hybrid polymers was increased with an increasing ratio of POSS moieties as the inorg. part. Because of the steric hindrance caused by the bulkiness of the POSS macromonomer, the number-average mol. weight of the hybrid polymers gradually decreased as the molar percentage of POSS in the feed increased. Their liquid crystallinities were very dependent on the POSS segments of the hybrid polymers behaving as hard, compact components. The hybrid polymer with 90 mol% LC moiety (Cube-LC90) showed liquid crystallinity, larger glass-transition temps., and better stability with respect to the LC homopolymer. The results of differential scanning calorimetry and optical polarizing microscopy showed that Cube-LC90 had a smectic-mesophase-like fine-grained texture.

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PENG 10/520763 07/11/2007Page 132
     35-4 (Chemistry of Synthetic High Polymers)
CC
     Section cross-reference(s): 75
ST
     liq cryst acrylic polymer polyhedral silsesquioxane side group
IT
     Steric hindrance
        (in liquid-crystalline acrylic copolymers with polyhedral silsesquioxane
side
        groups)
IT
     Liquid crystals, polymeric
        (liquid-crystalline acrylic copolymers with polyhedral silsesquioxane side
IT
     Glass transition temperature
     Thermal stability
        (of liquid-crystalline acrylic copolymers with polyhedral silsesquioxane
side
        groups)
IT
     Silsesquioxanes
    RL: PRP (Properties); RCT (Reactant); SPN (Synthetic preparation); PREP
     (Preparation); RACT (Reactant or reagent)
        (polyhedral, oligomeric; liquid-crystalline acrylic copolymers with
polyhedral
        silsesquioxane side groups)
TT
     14579-03-4, Cyclopentyltrichlorosilane
                                              38595-89-0, (3-...
     Acryloxypropyl) trichlorosilane
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (in monomer preparation; liquid-crystalline acrylic copolymers with
polyhedral
        silsesquioxane side groups)
IT
     183387-28-2P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (in monomer preparation; liquid-crystalline acrylic copolymers with
polyhedral
        silsesquioxane side groups)
IT
     117318-92-0P 387820-82-8P 387820-83-9P
     RL: PRP (Properties); SPN (Synthetic preparation); PREP
     (Preparation)
        (liquid-crystalline acrylic copolymers with polyhedral silsesquioxane side
        groups)
IT
     117318-91-9P 387820-81-7P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP
     (Preparation); RACT (Reactant or reagent)
        (monomer; liquid-crystalline acrylic copolymers with polyhedral
silsesquioxane
        side groups)
IT
     387820-82-8P 387820-83-9P
     RL: PRP (Properties); SPN (Synthetic preparation); PREP
     (Preparation)
        (liquid-crystalline acrylic copolymers with polyhedral silsesquioxane side
        groups)
RN
     387820-82-8 HCAPLUS
CN
     2-Propenoic acid, 2-methyl-, 6-[(4'-cyano[1,1'-biphenyl]-4-yl)oxy]hexyl
     ester, polymer with 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,15.17,13]o
     ctasiloxanyl)propyl 2-propenoate (9CI) (CA INDEX NAME)
     CM
          1
     CRN 387820-81-7
     CMF C41 H72 O14 Si8
```

PENG 10/520763 07/11/2007Page 133

CM 2

CRN 117318-91-9 CMF C23 H25 N O3

RN 387820-83-9 HCAPLUS

CN 2-Propenoic acid, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,15.17,13]oct asiloxanyl)propyl ester, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 387820-81-7 CMF C41 H72 O14 Si8

IT 387820-81-7P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP

(Preparation); RACT (Reactant or reagent)

(monomer; liquid-crystalline acrylic copolymers with polyhedral

silsesquioxane

side groups)

RN 387820-81-7 HCAPLUS

CN 2-Propenoic acid, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,15.17,13]oct asiloxanyl)propyl ester (9CI) (CA INDEX NAME)

RE.CNT 21 THERE ARE 21 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

KATHLEEN FULLER EIC1700 571/272-2505

```
ANSWER 19 OF 27 HCAPLUS COPYRIGHT 2007 ACS on STN
L37
AN
     2001:503820 HCAPLUS
DN
     136:74531
TI
     Evaluation of neat resins based on methacrylates modified with
     methacryl-POSS, as potential organic-inorganic hybrids for
     formulating dental restoratives
ΑU
     Gao, Feng; Tong, Yuhua; Schricker, Scott R.; Culbertson, Bill M.
     College of Dentistry, The Ohio State University, Columbus, OH, 43218-2357,
CS
     Polymers for Advanced Technologies (2001), 12(6), 355-360
SO
     CODEN: PADTE5; ISSN: 1042-7147
PB
     John Wiley & Sons Ltd.
DT
     Journal
     English
LA
AΒ
     Three series of neat resins, based on methacrylates or dimethacrylates
     modified with 1-[3-(methacryl)propyl]-3,5,7,9,11,13,15-
     heptacyclopentylpentacyclo[9.5.1.13,9.15,15.17,13]octasiloxane
     (Methacryl-POSS or POSS-MA), were evaluated. Incorporation of POSS-MA
     into the polymeric matrix was accomplished in three ways: (a) resins
     series I was formulated with one-pot copolymn. of a bis-GMA/diluent
     monomer with POSS-MA; (b) the copolymer of 2-hydroxylethyl methacrylate
     (HEMA) and POSS-MA, with a weight ratio of 1:1, was first made, then the
     copolymer hydroxyl groups were converted into methacrylate groups to get a
     macromer; finally the macromer was copolymd. with bis-GMA/diluents to
     formulate the second series of neat resins; (c) copolymer of POSS-MA and
     t-Bu methacrylate (t-BMA) or 2-ethylhexyl methacrylate (EHMA) were first
     made, then they were combined with a mixture of bis-GMA and the
     corresponding diluent to make the third kind of neat resins.
     POSS incorporated into the neat resin matrix was controlled at 5, 10 and
             Incorporating only 5 wt% substantially reduced the shrinkage of
     the prepared neat resins. The percentage of methacrylate-functionalized
     POSS monomer in the bis-GMA/diluent base, was compared to neat resins
     prepared without the POSS. The mech. properties of neat resin having POSS.
     were improved for approaches (b) and (c), but remained at the same level
     as the neat resins without POSS for approach (a). The results show that
     miscibility between the POSS component and the matrix, especially the diluent,
     plays a very important role in the improvement of the properties of the
     formulated thermosets.
     63-7 (Pharmaceuticals)
CC
     Section cross-reference(s): 35, 36
ST
     dental resin methacrylate siloxane
IT
     Compressive strength
     Contraction (mechanical)
        (evaluation of neat dental resins based on methacrylates modified with
        1-[3-(methacryl)propyl]-3,5,7,9,11,13,15-heptacyclopentylpentacyclo[9.5
        .1.13,9.15,15.17,13] octasiloxane as potential organic-inorg.
IT
     Polysiloxanes, biological studies
     RL: PRP (Properties); SPN (Synthetic preparation); THU (Therapeutic use);
     BIOL (Biological study); PREP (Preparation); USES (Uses)
        (methacrylate-; evaluation of neat dental resins based on methacrylates
        modified with 1-[3-(methacryl)propyl]-3,5,7,9,11,13,15-
        heptacyclopentylpentacyclo[9.5.1.13,9.15,15.17,13]octasiloxane as
        potential organic-inorg. hybrids)
IT
     Dental materials and appliances
        (resins; evaluation of neat dental resins based on methacrylates
```

modified with 1-[3-(methacryl)propyl]-3,5,7,9,11,13,15-

heptacyclopentylpentacyclo[9.5.1.13,9.15,15.17,13]octasiloxane as

potential organic-inorg. hybrids)

IT 279687-80-8P 279687-83-1P 279687-85-3P

384339-56-4P 384340-21-0P

RL: PRP (Properties); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(evaluation of neat dental resins based on methacrylates modified with 1-[3-(methacryl)propyl]-3,5,7,9,11,13,15-heptacyclopentylpentacyclo[9.5.1.13,9.15,15.17,13]octasiloxane as potential organic-inorg. hybrids)

IT 279687-80-8P 279687-83-1P 279687-85-3P

384339-56-4P 384340-21-0P

RL: PRP (Properties); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(evaluation of neat dental resins based on methacrylates modified with 1-[3-(methacryl)propyl]-3,5,7,9,11,13,15-heptacyclopentylpentacyclo[9.5.1.13,9.15,15.17,13]octasiloxane as potential organic-inorg.

hybrids)
RN 279687-80-8 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 1,6-hexanediyl ester, polymer with 3-(heptacyclopentylpentacyclo[9.5.1.13,?.15,15.17,13]octasiloxanyl)propyl 2-methyl-2-propenoate and (1-methylethylidene)bis[4,1-phenyleneoxy(2-hydroxy-3,1-propanediyl)] bis(2-methyl-2-propenoate) (9CI) (CA INDEX NAME)

CM 1

CRN 169391-91-7 CMF C42 H74 O14 Si8

CM 2

CRN 6606-59-3 CMF C14 H22 O4

CRN 1565-94-2 CMF C29 H36 O8

PAGE 1-A

PAGE 1-B

$$\begin{array}{c|c} \text{O} & \text{CH}_2 \\ || & || \\ -\text{C-C-Me} \end{array}$$

RN 279687-83-1 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, (1-methylethylidene)bis[4,1-phenyleneoxy(2-hydroxy-3,1-propanediyl)] ester, polymer with 1,1-dimethylethyl 2-methyl-2-propenoate and 3-(heptacyclopentylpentacyclo[9.5.1.13,?.15,15.1 7,13]octasiloxanyl)propyl 2-methyl-2-propenoate (9CI) (CA INDEX NAME)

CM 1

CRN 169391-91-7 CMF C42 H74 O14 Si8

PENG 10/520763 07/11/2007Page 138

CM · 2

CRN 1565-94-2 CMF C29 H36 O8

PAGE 1-A

PAGE 1-B

$$\begin{array}{c|c} \text{O} & \text{CH}_2 \\ || & || \\ -\text{C-C-Me} \end{array}$$

CM 3

CRN 585-07-9 CMF C8 H14 O2

$$\begin{array}{c|c} \text{O} & \text{CH}_2 \\ \parallel & \parallel \\ \text{t-BuO-C-C-Me} \end{array}$$

RN 279687-85-3 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, (1-methylethylidene)bis[4,1-phenyleneoxy(2-hydroxy-3,1-propanediyl)] ester, polymer with 2-ethylhexyl 2-methyl-2-propenoate and 3-(heptacyclopentylpentacyclo[9.5.1.13,?.15,15.17,13]octasiloxanyl)propyl 2-methyl-2-propenoate (9CI) (CA INDEX NAME)

CM 1

CRN 169391-91-7 CMF C42 H74 O14 Si8

CM 2

CRN 1565-94-2 CMF C29 H36 O8

CRN 688-84-6 CMF C12 H22 O2

$$\begin{array}{c|c} & \text{O} & \text{CH}_2 \\ \parallel & \parallel \\ \text{CH}_2-\text{O-C-C-Me} \\ \parallel \\ \text{Et-CH-Bu-n} \end{array}$$

RN 384339-56-4 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 1,6-hexanediyl ester, polymer with 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,15.17,13]octasiloxanyl)propyl 2-methyl-2-propenoate and α,α' -[(1-methylethylidene)di-4,1-phenylene]bis[ω -[(2-methyl-1-oxo-2-propenyl)oxy]poly(oxy-1,2-ethanediyl)] (9CI) (CA INDEX NAME)

CM 1

CRN 169391-91-7 CMF C42 H74 O14 Si8

CM 2

CRN 41637-38-1 CMF (C2 H4 O)n (C2 H4 O)n C23 H24 O4 CCI PMS

PAGE 1-A

PAGE 1-B

$$\begin{array}{c|c} & O & CH_2 \\ \hline & & & \\ & & \\ \hline & & \\ & & \\ \end{array}$$

CM 3

CRN 6606-59-3 CMF C14 H22 O4

RN 384340-21-0 HCAPLUS

2-Propenoic acid, 2-methyl-, 1,2-ethanediylbis(oxy-2,1-ethanediyl) ester, polymer with 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,15.17,13]octasilo xanyl)propyl 2-methyl-2-propenoate polymer with 2-hydroxyethyl 2-methyl-2-propenoate 2-methyl-2-propenoate, and (1-methylethylidene)bis[4,1-phenyleneoxy(2-hydroxy-3,1-propanediyl)] bis(2-methyl-2-propenoate) (9CI) (CA INDEX NAME)

CM 1

CRN 1565-94-2 CMF C29 H36 O8

PAGE 1-B

CM 2

CRN 109-16-0 CMF C14 H22 O6

CM 3

CRN 384340-20-9

CMF (C42 H74 O14 Si8 . C6 H10 O3)x . x C4 H6 O2

CM 4

CRN 79-41-4 CMF C4 H6 O2

$$^{{
m CH}_2}_{||}$$
 Me- C- $^{{
m CO}_2}$ H

CM 5

CRN 384340-19-6

CMF (C42 H74 O14 Si8 . C6 H10 O3)x

CCI PMS

CM 6

CRN 169391-91-7 CMF C42 H74 O14 Si8

KATHLEEN FULLER EIC1700 571/272-2505

CRN 868-77-9 CMF C6 H10 O3

RE.CNT 6 THERÉ ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L37 ANSWER 20 OF 27 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2000:428857 HCAPLUS

DN 133:177504

TI Hydrosilylation of Allyl Alcohol with [HSiMe2OSiO1.5]8:
Octa(3-hydroxypropyldimethylsiloxy)octasilsesquioxane and Its
Octamethacrylate Derivative as Potential Precursors to Hybrid
Nanocomposites

AU Zhang, Chunxin; Laine, Richard M.

CS Departments of Chemistry and Materials Science and Engineering and the Macromolecular Science and Engineering Center, University of Michigan, Ann Arbor, MI, 48109-2136, USA

SO Journal of the American Chemical Society (2000), 122(29), 6979-6988 CODEN: JACSAT; ISSN: 0002-7863

PB American Chemical Society

DT Journal

LA English

AB Octakis(3-hydroxypropyldimethylsiloxy)octasilsesquioxane (OHPS) was synthesized by direct hydrosilylation of allyl alc. with octakis(dimethylsiloxy)octasilsesquioxane, (HSiMe2O)Si8O12 (Q8M8H), using platinum divinyltetramethyldisiloxane [Pt(dvs)] as catalyst.

10/520763 07/11/2007Page 144 Surprisingly, C-hydrosilylation occurs in preference to O-silylation. Hydrosilylation of trimethylsiloxy-2-propene with Q8M8H, followed by desilylation also gives pure OHPS. The OHPS reacts with methacryloyl chloride to give octakis(3-methacryloxypropyldimethylsiloxy)octasilsesquioxane (OMPS), a thermal and UV/vis curable precursor to organic/inorg. nanocomposites. Direct hydrosilylation of Q8M8H with 2-allyloxyethanol also proceeds primarily via C- rather than O-silylation. In contrast, compds. such as 1,3,5,7tetramethylcyclotetrasiloxane (D4H), 1,1,3,3-tetramethyldisiloxane (TMDS) or terminal Si-H functionalized poly(dimethylsiloxane) (PDMS-H, MW = 400), give significant amts. of O-silylation along with C-silylation. Initial catalyst concentration studies suggest that the catalytic cycle requires the intermediacy of Pt cluster complexes in contrast to recent studies on the mechanism of hydrosilylation which suggest monometallic complex catalysis. 35-2 (Chemistry of Synthetic High Polymers) Section cross-reference(s): 37, 57 allyl alc hydrosilylation methylsiloxysilsesquioxane hydroxypropyldimethylsiloxy silsesquioxane prepn; silsesquioxane prepn platinum vinylmethyldisiloxane hydrosilylation catalyst; hybrid nanocomposite precursor silsesquioxane prepn hydrosilylation Silylation (C-hydrosilylation, O-silylation, C-silylation; hydrosilylation of allyl alc. with octasilsesquioxane and preparation of octamethacrylate derivative precursors to hybrid nanocomposites) Addition reaction kinetics Addition reaction kinetics (hydrosilylation kinetics; hydrosilylation of allyl alc. with octasilsesquioxane and preparation of octamethacrylate derivative precursors to hybrid nanocomposites) Hybrid organic-inorganic materials Hydrosilylation catalysts Solvent effect (hydrosilylation of allyl alc. with octasilsesquioxane and preparation of octamethacrylate derivative precursors to hybrid nanocomposites) RL: SPN (Synthetic preparation); PREP (Preparation) (hydrosilylation of allyl alc. with octasilsesquioxane and preparation of octamethacrylate derivative precursors to hybrid nanocomposites) Polysiloxanes, reactions RL: RCT (Reactant); RACT (Reactant or reagent) (hydrosilylation reactant; hydrosilylation of allyl alc. with octasilsesquioxane and preparation of octamethacrylate derivative hybrid nanocomposites) Hydrosilylation Hydrosilylation (kinetics; hydrosilylation of allyl alc. with octasilsesquioxane and preparation of octamethacrylate derivative precursors to hybrid

ΙT

CC

IT

IT

IT

IT

IT

nanocomposites)

IT 16941-12-1, Hexachloroplatinic acid

RL: CAT (Catalyst use); USES (Uses)

(hydrosilylation catalyst; hydrosilylation of allyl alc. with octasilsesquioxane and preparation of octamethacrylate derivative precursors to

hybrid nanocomposites)

7440-06-4DP, Platinum, cyclopentadiene complexes, preparation RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent) (hydrosilylation catalyst; hydrosilylation of allyl alc. with

octasilsesquioxane and preparation of octamethacrylate derivative precursors to hybrid nanocomposites) IT 107-18-6, 2-Propen-1-ol, reactions 107-19-7, Propargyl alcohol 111-45-5, 2-Allyloxyethanol 920-46-7 18146-00-4, Allyloxytrimethylsilane 125756-69-6, Octakis(dimethylsiloxy)octasilsesqu ioxane RL: RCT (Reactant); RACT (Reactant or reagent) (hydrosilylation of allyl alc. with octasilsesquioxane and preparation of octamethacrylate derivative precursors to hybrid nanocomposites) IT 2370-88-9, Tetramethylcyclotetrasiloxane 9016-00-6, Di-Me siloxane, SRU 30110-74-8, Tetramethyldisiloxane 31900-57-9, Dimethylsilanediol homopolymer RL: RCT (Reactant); RACT (Reactant or reagent) (hydrosilylation reactant; hydrosilylation of allyl alc. with octasilsesquioxane and preparation of octamethacrylate derivative precursors to hybrid nanocomposites) 75-09-2, Dichloromethane, uses 108-88-3, Toluene, uses 109-99-9, uses 142-82-5, Heptane, uses RL: NUU (Other use, unclassified); USES (Uses) (hydrosilylation solvent; hydrosilylation of allyl alc. with octasilsesquioxane and preparation of octamethacrylate derivative precursors to hybrid nanocomposites) IT 288290-32-4P, Octakis (3-hydroxypropyldimethylsiloxy) octasilsesquioxane RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent) (intermediate and precursor; hydrosilylation of allyl alc. with octasilsesquioxane and preparation of octamethacrylate derivative precursors to hybrid nanocomposites) TT 288290-33-5P, Octakis[(3-trimethylsiloxypropyl)dimethylsiloxy]silsesquioxa RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent) (intermediate; hydrosilylation of allyl alc. with octasilsesquioxane and preparation of octamethacrylate derivative precursors to hybrid nanocomposites) IT 288290-34-6P, Octakis(3-methacryloxypropyldimethylsiloxy)octasilse squioxane RL: SPN (Synthetic preparation); PREP (Preparation) (precursor; hydrosilylation of allyl alc. with octasilsesquioxane and preparation of octamethacrylate derivative precursors to hybrid nanocomposites) IT 288290-34-6P, Octakis (3-methacryloxypropyldimethylsiloxy) octasilse squioxane RL: SPN (Synthetic preparation); PREP (Preparation) (precursor; hydrosilylation of allyl alc. with octasilsesquioxane and

- nanocomposites)
 RN 288290-34-6 HCAPLUS
- CN 2-Propenoic acid, 2-methyl-, pentacyclo[9.5.1.13,9.15,15.17,13]octasiloxan e-1,3,5,7,9,11,13,15-octayloctakis[oxy(dimethylsilylene)-3,1-propanediyl] ester (9CI) (CA INDEX NAME)

preparation of octamethacrylate derivative precursors to hybrid

Me

CH₂ O

CH₂

PAGE 3-B

$$-$$
 (CH₂)₃-O-C-C-Me

RE.CNT 72 THERE ARE 72 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L37 ANSWER 21 OF 27 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2000:208313 HCAPLUS

DN 132:335664

TI Synthesis and microstructural characterization of POSS-based triblock copolymers prepared using atom transfer radical polymerization

AU Mather, Patrick T.; Chun, Seung B.; Pyun, Jeffrey; Matyjaszewski, Krzysztof; Jeon, Hong G.

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CS
     Polymer Program, Institute of Materials Science, University of
     Connecticut, Storrs, CT, 06269-3136, USA
SO
     Polymer Preprints (American Chemical Society, Division of Polymer
     Chemistry) (2000), 41(1), 582-583
     CODEN: ACPPAY; ISSN: 0032-3934
PR
     American Chemical Society, Division of Polymer Chemistry
DT
     Journal
LA
     English
AB
     POSS-based hybrid polymers have provided an approach toward
     enhancing the properties of existing polymer systems with little
     modification to polymerization methods. The present challenge is to extend
this
     paradigm to more complex macromol. architectures and compns., namely block
     copolymers. Block copolymers incorporating polyhedral oligomeric
     silsesquioxane (POSS) monomers using atom transfer radical polymerization
     are made and characterized. Triblock copolymers have been synthesized
     starting from polyacrylate macroinitiators (p(Bu acrylate)) and chain
     extending with methacrylate POSS monomers. Mol. characterization of these
     materials was conducted by SEC and 1H NMR, while microstructural
     characterization was performed using transmission electron microscopy
     (TEM), small angle x-ray scattering (SAXS), and wide-angle x-ray
     scattering (WAXS). The ordered microstructures evolve in terms of the
     interactions between the POSS outer blocks and the acrylate inner block.
     39-4 (Synthetic Elastomers and Natural Rubber)
CC
ST
     methacryloyl silsesquioxane butyl acrylate block copolymer
IT
     Glass transition temperature
       Hybrid organic-inorganic materials
        (acrylic-silsesquioxane triblock copolymers prepared using atom transfer
        radical polymerization)
IT
    Acrylic rubber
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (block; acrylic-silsesquioxane triblock copolymers prepared using atom
        transfer radical polymerization)
IT
     Polymerization
        (block; synthesis and microstructural characterization of POSS-based
        triblock copolymers prepared using atom transfer radical polymerization)
IT
        (radical, atom transfer; synthesis and microstructural characterization
        of POSS-based triblock copolymers prepared using)
IT
     255872-36-7P 841235-76-5P
    RL: SPN (Synthetic preparation); PREP (Preparation)
        (rubber; acrylic-silsesquioxane triblock copolymers prepared using atom
        transfer radical polymerization)
IT
     255872-36-7P 841235-76-5P
    RL: SPN (Synthetic preparation); PREP (Preparation)
        (rubber; acrylic-silsesquioxane triblock copolymers prepared using atom
       transfer radical polymerization)
RN
    255872-36-7 HCAPLUS
CN
     2-Propenoic acid, 2-methyl-, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,1
     5.17,13]octasiloxanyl)propyl ester, polymer with butyl 2-propenoate, block
     (9CI)
            (CA INDEX NAME)
```

CRN 141-32-2 CMF C7 H12 O2

$$\begin{array}{c} \text{O} \\ \parallel \\ \text{n-BuO-C-CH} \end{array}$$

RN 841235-76-5 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,1 5.17,13]octasiloxanyl)propyl ester, polymer with butyl 2-propenoate, triblock (9CI) (CA INDEX NAME)

CM 1

CRN 141-32-2 CMF C7 H12 O2

RE.CNT 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L37 ANSWER 22 OF 27 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 2000:208312 HCAPLUS

DN 133:79275

TI Evaluation of multi-methacrylates copolymerized with methacryl-POSS for potential organic-inorganic hybrid dental restorative materials

AU Gao, Feng; Culbertson, Bill. M.; Tong, Yuhua; Schricker, Scott R.

CS Dentistry College, The Ohio State University, Columbus, OH, 43210-1241, USA

SO Polymer Preprints (American Chemical Society, Division of Polymer Chemistry) (2000), 41(1), 580-581
CODEN: ACPPAY; ISSN: 0032-3934

PB American Chemical Society, Division of Polymer Chemistry

DT Journal

LA English

AB The shrinkage of methacrylate-based dental resins prepared can be reduced efficiently by incorporating 5 weight% of methacrylate-functionalized Polyhedral Oligomeric Silsesquioxane (POSS) macromonomer into a mixture of oligomeric dimethacrylate/diluent mixture The mech. properties of the unfilled resins can also be improved.

CC 63-7 (Pharmaceuticals)

ST dental restoration methacrylate polyhedral oligomer silsesquioxane

IT Silsesquioxanes

RL: PRP (Properties); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(evaluation of multi-methacrylates copolymd. with methacryl-polyhedral oligomeric silsesquioxane for potential organic-inorg. hybrid dental restorative materials)

IT Dental materials and appliances

(resins; evaluation of multi-methacrylates copolymd. with methacryl-polyhedral oligomeric silsesquioxane for potential organic-inorg. hybrid dental restorative materials)

IT 279687-80-8P 279687-83-1P 279687-85-3P

RL: PRP (Properties); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (evaluation of multi-methacrylates copolymd. with methacryl-polyhedral

oligomeric silsesquioxane for potential organic-inorg. hybrid dental restorative materials)

279687-80-8P 279687-83-1P 279687-85-3P

RL: PRP (Properties); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(evaluation of multi-methacrylates copolymd. with methacryl-polyhedral oligomeric silsesquioxane for potential organic-inorg. hybrid dental restorative materials)

RN 279687-80-8 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 1,6-hexanediyl ester, polymer with 3-(heptacyclopentylpentacyclo[9.5.1.13,?.15,15.17,13]octasiloxanyl)propyl 2-methyl-2-propenoate and (1-methylethylidene)bis[4,1-phenyleneoxy(2-hydroxy-3,1-propanediyl)] bis(2-methyl-2-propenoate) (9CI) (CA INDEX NAME)

CM 1

IT

CRN 6606-59-3 CMF C14 H22 O4

CM 3

CRN 1565-94-2 CMF C29 H36 O8

PAGE 1-A

PAGE 1-B

$$\begin{array}{c|c} \text{O} & \text{CH}_2 \\ || & || \\ -\text{C-C-Me} \end{array}$$

RN 279687-83-1 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, (1-methylethylidene)bis[4,1-phenyleneoxy(2-hydroxy-3,1-propanediyl)] ester, polymer with 1,1-dimethylethyl 2-methyl-2-propenoate and 3-(heptacyclopentylpentacyclo[9.5.1.13,?.15,15.1 7,13]octasiloxanyl)propyl 2-methyl-2-propenoate (9CI) (CA INDEX NAME)

CM 1

CM 2

CRN 1565-94-2 CMF C29 H36 O8

PAGE 1-A

$$H_{2}^{C}$$
 O OH OH Me-C-C-O-CH₂-CH-CH₂-O Me O-CH₂-CH-CH₂-O Me

PAGE 1-B

CM :

CRN 585-07-9 CMF C8 H14 O2

$$\begin{array}{c|c} & \text{O} & \text{CH}_2 \\ & || & || \\ \text{t-BuO-} & \text{C--C-Me} \end{array}$$

RN 279687-85-3 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, (1-methylethylidene)bis[4,1-phenyleneoxy(2-hydroxy-3,1-propanediyl)] ester, polymer with 2-ethylhexyl 2-methyl-2-propenoate and 3-(heptacyclopentylpentacyclo[9.5.1.13,?.15,15.17,13]octasiloxanyl)propyl 2-methyl-2-propenoate (9CI) (CA INDEX NAME)

CM 1

CRN 169391-91-7 CMF C42 H74 O14 Si8

CM 2

CRN 1565-94-2 CMF C29 H36 O8

$$\begin{array}{c|c} & \text{O} & \text{CH}_2 \\ & || & || \\ & -\text{C--C-Me} \end{array}$$

CRN 688-84-6 CMF C12 H22 O2

RE.CNT 7

THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L37 ANSWER 23 OF 27 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 1999:780676 HCAPLUS

DN 132:108370

TI Synthesis of hybrid polymers Using atom transfer radical polymerization: Homopolymers and Block Copolymers from polyhedral oligomeric silsesquioxane monomers

AU Pyun, Jeffrey; Matyjaszewski, Krzysztof

CS Center for Macromolecular Engineering Department of Chemistry, Carnegie Mellon University, Pittsburgh, PA, 15213, USA

SO Macromolecules (2000), 33(1), 217-220 CODEN: MAMOBX; ISSN: 0024-9297

American Chemical Society

DT Journal

PB

LA English

AB Homopolymers, triblock copolymers (Bu acrylate comonomer), and star-block copolymers (Me acrylate comonomer) of 3-(3,5,7,9,11,13,15-heptacyclopentyl-pentacyclo[9.5.1.1.3,91.5,1517,13]octasiloxane-1-yl)propyl methacrylate (MA-POSS) have been prepared

CC 35-4 (Chemistry of Synthetic High Polymers)

ST methacrylate functional silsesquioxane atom transfer radical polymn; block methacrylate functional silsesquioxane; star block methacrylate functional silsesquioxane; butyl acrylate methacrylate functional silsesquioxane copolymer; methyl acrylate methacrylate functional silsesquioxane copolymer

IT Polymers, preparation

RL: SPN (Synthetic preparation); PREP (Preparation) (block, siloxanyl star-block; synthesis of hybrid polymers using atom transfer radical polymerization of polyhedral oligomeric silsesquioxane monomers)

IT Polymerization

(radical, atom-transfer; synthesis of hybrid polymers using atom transfer radical polymerization of polyhedral oligomeric silsesquioxane monomers)

radical polymerization of polyhedral oligomeric silsesquioxane monomers)

IT 169699-57-4P

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation and characterization of; synthesis of hybrid polymers using atom transfer radical polymerization of polyhedral oligomeric silsesquioxane monomers)

IT 255872-37-8P

RL: SPN (Synthetic preparation); PREP (Preparation)
 (star-block, 3-arm, preparation and characterization of; synthesis of
 hybrid polymers using atom transfer radical polymerization of
 polyhedral oligomeric silsesquioxane monomers)

IT 255872-36-7P 841235-76-5P

RL: SPN (Synthetic preparation); PREP (Preparation)
(triblock, preparation and characterization of; synthesis of hybrid
polymers using atom transfer radical polymerization of polyhedral oligomeric
silsesquioxane monomers)

IT 169699-57-4P

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation and characterization of; synthesis of hybrid polymers using atom transfer radical polymerization of polyhedral oligomeric silsesquioxane monomers)

RN 169699-57-4 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,1 5.17,13]octasiloxanyl)propyl ester, homopolymer (9CI) (CA INDEX NAME)

· CM 1

CRN 169391-91-7 CMF C42 H74 O14 Si8

IT 255872-37-8P

RL: SPN (Synthetic preparation); PREP (Preparation) (star-block, 3-arm, preparation and characterization of; synthesis of hybrid polymers using atom transfer radical polymerization of polyhedral oligomeric silsesquioxane monomers)

RN 255872-37-8 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,1 5.17,13]octasiloxanyl)propyl ester, polymer with methyl 2-propenoate, block (9CI) (CA INDEX NAME)

CM 1

CRN 169391-91-7 CMF C42 H74 O14 Si8

CM 2

CRN 96-33-3 CMF C4 H6 O2

 $\begin{array}{c} \text{O} \\ || \\ \text{MeO-C-CH-----} \text{CH}_2 \end{array}$

IT 255872-36-7P 841235-76-5P

RL: SPN (Synthetic preparation); PREP (Preparation)
(triblock, preparation and characterization of; synthesis of hybrid
polymers using atom transfer radical polymerization of polyhedral oligomeric
silsesquioxane monomers)

RN 255872-36-7 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,1 5.17,13]octasiloxanyl)propyl ester, polymer with butyl 2-propenoate, block (9CI) (CA INDEX NAME)

CM 1

CRN 141-32-2 CMF C7 H12 O2

RN 841235-76-5 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,1 5.17,13]octasiloxanyl)propyl ester, polymer with butyl 2-propenoate, triblock (9CI) (CA INDEX NAME)

CM 1

CRN 141-32-2 CMF C7 H12 O2

RE.CNT 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CYTATIONS AVAILABLE IN THE RE FORMAT

L37 ANSWER 24 OF 27 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 1999:558896 HCAPLUS

DN 132:166622

TI Synthesis of organic/inorganic hybrid materials from polysiloxane precursors using atom transfer radical polymerization

AU Pyun, Jeffrey; Miller, Peter J.; Kickelbick, Guido; Matyjaszewski, Krzysztof; Schwab, Joseph; Lichtenhan, Joseph

CS Center for Macromolecular Engineering, Dept. of Chemistry, Carnegie Mellon University, Pittsburgh, PA, 15213, USA

SO Polymer Preprints (American Chemical Society, Division of Polymer Chemistry) (1999), 40(2), 454-455
CODEN: ACPPAY; ISSN: 0032-3934

PB American Chemical Society, Division of Polymer Chemistry

DT Journal

LA English

AB The synthesis of organic/inorg. hybrid materials from polyhedral oligomeric silsesquioxanes (POSS) using atom transfer radical polymerization (ATRP) was carried out. From POSS monomers containing either methacryloyl, or styryl groups, well-defined polymers of various compns. and topologies were prepared The composition of these materials was controlled and well-defined

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homopolymers, random and block copolymers of POSS-containing monomers were synthesized. The use of macroinitiators of various topologies allowed for the synthesis of ABA-triblock and star-block copolymers, using ATRP. 35-4 (Chemistry of Synthetic High Polymers) Section cross-reference(s): 37, 57 silsesquioxane polyhedral oligomer atom transfer radical polymn; methacryloyl styryl silsesquioxane ATRP polysiloxane precursor hybrid; hybrid material prepn polyhedral polysiloxane precursor Silsesquioxanes RL: SPN (Synthetic preparation); PREP (Preparation) (methacryloyl- and styryl-; preparation of methacryloyl- and styryl-silsesquioxane homopolymers and block copolymers and star copolymers by ATRP as precursors for organic/inorg. hybrid materials) Hybrid organic-inorganic materials Molecular topology (preparation of methacryloyl- and styryl-silsesquioxane homopolymers and block copolymers and star copolymers by ATRP as precursors for organic/inorg. hybrid materials) Polysiloxanes, preparation RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of methacryloyl- and styryl-silsesquioxane homopolymers and block copolymers and star copolymers by ATRP as precursors for organic/inorg. hybrid materials) Polymerization (radical, atom transfer; preparation of methacryloyl- and styryl-silsesquioxane homopolymers and block copolymers and star copolymers by ATRP as precursors for organic/inorg. hybrid materials) Polymers, preparation RL: SPN (Synthetic preparation); PREP (Preparation) (star-branched; preparation of methacryloyl- and styryl-silsesquioxane homopolymers and block copolymers and star copolymers by ATRP as precursors for organic/inorg. hybrid materials) 255872-36-7P RL: SPN (Synthetic preparation); PREP (Preparation) (block; preparation of methacryloyl- and styryl-silsesquioxane homopolymers and block copolymers and star copolymers by ATRP as precursors for organic/inorg. hybrid materials) 600-00-0, Ethyl-2-bromoisobutyrate RL: CAT (Catalyst use); USES (Uses) (initiator; preparation of methacryloyl- and styryl-silsesquioxane homopolymers and block copolymers and star copolymers by ATRP as precursors for organic/inorg. hybrid materials) 3030-47-5, PMDETA RL: NUU (Other use, unclassified); USES (Uses) (preparation of methacryloyl- and styryl-silsesquioxane homopolymers and block copolymers and star copolymers by ATRP as precursors for organic/inorg. hybrid materials) 169699-57-4P

IT

RL: PNU (Preparation, unclassified); PREP (Preparation) (preparation of methacryloyl- and styryl-silsesquioxane homopolymers and block copolymers and star copolymers by ATRP as precursors for organic/inorg. hybrid materials)

183200-99-9P IT 169391-91-7P

> RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of methacryloyl- and styryl-silsesquioxane homopolymers and block copolymers and star copolymers by ATRP as precursors for organic/inorg. hybrid materials)

IT 7758-89-6, Cuprous chloride

RL: CAT (Catalyst use); USES (Uses)

(radical polymerization catalyst; preparation of methacryloyl- and styryl-silsesquioxane homopolymers and block copolymers and star copolymers by ATRP as precursors for organic/inorg. hybrid materials)

IT 255872-37-8P

RL: PNU (Preparation, unclassified); PREP (Preparation)
(star block; preparation of methacryloyl- and styryl-silsesquioxane homopolymers and block copolymers and star copolymers by ATRP as precursors for organic/inorg. hybrid materials)

IT 255872-36-7P

RL: SPN (Synthetic preparation); PREP (Preparation)
(block; preparation of methacryloyl- and styryl-silsesquioxane homopolymers and block copolymers and star copolymers by ATRP as precursors for organic/inorg. hybrid materials)

RN 255872-36-7 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,1 5.17,13]octasiloxanyl)propyl ester, polymer with butyl 2-propenoate, block (9CI) (CA INDEX NAME)

CM 1

CRN 169391-91-7 CMF C42 H74 O14 Si8

CM 2

CRN 141-32-2 CMF C7 H12 O2

IT 169699-57-4P

RL: PNU (Preparation, unclassified); PREP (Preparation)
(preparation of methacryloyl- and styryl-silsesquioxane homopolymers and block copolymers and star copolymers by ATRP as precursors for organic/inorg. hybrid materials)

RN 169699-57-4 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,1 5.17,13]octasiloxanyl)propyl ester, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 169391-91-7 CMF C42 H74 O14 Si8

IT 169391-91-7P

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of methacryloyl- and styryl-silsesquioxane homopolymers and block copolymers and star copolymers by ATRP as precursors for organic/inorg. hybrid materials)

RN 169391-91-7 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,1 5.17,13]octasiloxanyl)propyl ester (9CI) (CA INDEX NAME)

IT 255872-37-8P

RL: PNU (Preparation, unclassified); PREP (Preparation)
(star block; preparation of methacryloyl- and styryl-silsesquioxane homopolymers and block copolymers and star copolymers by ATRP as precursors for organic/inorg. hybrid materials)

RN 255872-37-8 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,1 5.17,13]octasiloxanyl)propyl ester, polymer with methyl 2-propenoate, block (9CI) (CA INDEX NAME)

CM 1

CRN 96-33-3 CMF C4 H6 O2

RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L37 ANSWER 25 OF 27 CHCAPLUS COPYRIGHT 2007 ACS on STN

AN 1997:224374 HCAPLUS

DN 126:264553

TI Investigations into structure/property relationships for polyhedral oligomeric silsesquioxane (POSS) based methacrylate polymers

AU Schwab, Joseph J.; Lichtenhan, Joseph D.; Kevin, P.; Chaffee, P.; Carr, Michael J.; Bolf, Alan G.

CS Hughes STX, Phillips Lab., Edwards AFB, CA, 93524, USA

Polymer Preprints (American Chemical Society, Division of Polymer Chemistry) (1997), 38(1), 518-519
CODEN: ACPPAY; ISSN: 0032-3934

PB American Chemical Society, Division of Polymer Chemistry

DT Journal

LA English

AB Well-defined, thermoplastic **hybrid** polyhedral oligomeric silsesquioxane-based methacrylate polymers and their monomers were prepared The thermomech. data, oxygen permeability, and microstructure of the polymers were presented.

CC 36-5 (Physical Properties of Synthetic High Polymers)

ST polyhedral oligomeric silsesquioxane polymethacrylate synthesis microstructure; thermomech data silsesquioxane based polymethacrylate;

methacrylate polymer silsesquioxane based oxygen permeability

IT Permeability

(gas, oxygen; microstructure, thermomech. data, and oxygen permeability of polyhedral oligomeric silsesquioxane-based methacrylate polymers)

IT 164356-44-9

RL: RCT (Reactant); RACT (Reactant or reagent)

(Kin preparation of polyhedral oligomeric silsesquioxane-based methacrylate polymers)

IT 79-41-4, Methacrylic acid, reactions 4115-83-7 7351-61-3,

3-Trichlorosilylpropyl methacrylate 135225-24-0

RL: RCT (Reactant); RACT (Reactant or reagent)

(in preparation of polyhedral oligomeric silsesquioxane-based methacrylate polymers)

IT 169391-90-6P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP

(Preparation); RACT (Reactant or reagent)

(in preparation of polyhedral oligomeric silsesquioxane-based methacrylate polymers)

IT 169391-91-7P

RL: SPN (Synthetic preparation); PREP (Preparation)

(in preparation of polyhedral oligomeric silsesquioxane-based methacrylate polymers)

IT 9011-14-7, PMMA

RL: POF (Polymer in formulation); PRP (Properties); USES (Uses) (thermomech. data and oxygen permeability of blends of polyhedral

oligomeric silsesquioxane-based methacrylate polymers with PMMA)

IT 169391-90-6P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP

(Preparation); RACT (Reactant or reagent)

(in preparation of polyhedral oligomeric silsesquioxane-based methacrylate polymers)

RN 169391-90-6 HCAPLUS

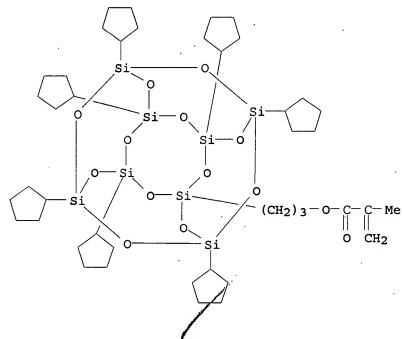
CN 2-Propenoic acid, 2-methyl-, 3-(heptacyclohexylpentacyclo[9.5.1.13,9.15,15.17,13]octasiloxanyl)propyl ester (9CI) (CA INDEX NAME)

IT 169391-91-7P

RL: SPN (Synthetic preparation); PREP (Preparation)
(in preparation of polyhedral oligomeric silsesquioxane-based methacrylate polymers)

RN 169391-91-7 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,1 5.17,13]octasiloxanyl)propyl ester (9CI) (CA INDEX NAME)



L37 ANSWER 26 OF 27 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 1995:983045 HCAPLUS

DN 124:30463

TI A One-Step Method for the Synthesis of a Vinyl-Containing Silsesquioxane and Other Organolithic Macromolecular Precursors

AU Yuchs, Steven E.; Carrado, Kathleen A.

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SO Inorganic Chemistry (1996), 35(1), 261-2

CODEN: INOCAJ; ISSN: 0020-1669

PB American Chemical Society

DT Journal

LA English

GI

AB A unique 1-step synthesis method for the introduction of a vinyl-containing functional group into an octasilsequioxane, [Si8012] (OR)8 (I) was developed. The target mol., I, R = CH2=Ch(Me) (Me)SiO, is synthesized using Me4N silicate and chlorodimethylvinylsilane. The mixture is stirred in a mixed organic phase for 1 h, and then hydrolyzed to complete the formation of the siloxane. The target mol., along with the Br and malonate derivs., were synthesized and characterized by a variety of traditional techniques. These silsesquioxane-based derivs. are valuable compds. which can be used to form hybrid polymers of silicic acids and new ceramic materials. This synthesis method for a vinyl containing silsesquioxane provides a very simple, high yield starting material used in the production of organolithic materials with varied properties.

CC 35-2 (Chemistry of Synthetic High Polymers)
 Section cross-reference(s): 57

ST vinyl silsesquioxane one step macromol precursor

IT Silsesquioxanes

RL: SPN (Synthetic preparation); PREP (Preparation)

(one-step method for synthesis of vinyl-containing silsesquioxane and other organolithic macromol. precursors)

Ι

IT 126503-69-3P, Octakis (vinyldimethylsiloxy) octasilsesquioxane 171979-93-4P 171979-94-5P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP

(Preparation); RACT (Reactant or reagent)

(one-step method for synthesis of vinyl-containing silsesquioxane and other organolithic macromol. precursors)

IT 1719-58-0, Chlorodimethylvinylsilane 53116-81-7, Tetramethylammonium silicate

RL: RCT (Reactant); RACT (Reactant or reagent)

(reactant; one-step method for synthesis of vinyl-containing silsesquioxane and other organolithic macromol. precursors)

IT 171979-94-5P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP

(Preparation); RACT (Reactant or reagent)

(one-step method for synthesis of vinyl-containing silsesquioxane and other organolithic macromol. precursors)

RN 171979-94-5 HCAPLUS

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

L37 ANSWER 27 OF 27 HCAPLUS COPYRIGHT 2007 ACS on STN

AN 1995:894380 HCAPLUS

DN 123:286932

TI Linear Hybrid Polymer Building Blocks: Methacrylate-

Functionalized Polyhedral Oligomeric Silsesquioxane Monomers and Polymers

AU Lichtenhan, Joseph D.; Otonari, Yoshiko A.; Carr, Michael J.

CS Propulsion Directorate, Phillips Laboratory, Edwards Air Force Base, CA, 93524, USA

SO Macromolecules (1995), 28(24), 8435-7

CODEN: MAMOBX; ISSN: 0024-9297

PB American Chemical Society

DT Journal

LA English

AB A new class of methacrylic monomers and their corresponding linear polymers have been developed. POSS monomers of the formula R7Si8O12(CH2)3OC(O)C(CH3):CH2 were synthesized by corner capping the POSS trisilanol cages R7Si7O9(OH)3 (where R = c-C6H11, c-C5H9) with methacrylate containing trichlorosilanes. The utility of these reagents for the preparation of linear methacrylate-based polymers containing the octameric silsesquioxane cage structure as a pendant group was demonstrated. Homo and copolymers of these systems are amorphous in nature and do not show any thermal transitions below their 388° decomposition temps. The thermal behavior of these systems is attributed to the dominant presence of the pendant silsesquioxane cages.

CC 35-4 (Chemistry of Synthetic High Polymers)

ST methacrylate functionalized silsesquioxane synthesis radical polymn

IT Polymer morphology

(of methacrylate-functionalized polyhedral oligomeric silsesquioxane polymers)

C42 H74 O14 Si8

PENG

CMF

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CRN 169391-90-6 CMF C49 H88 O14 Si8

IT 169699-56-3P 169699-57-4P

RL: PRP (Properties); SPN (Synthetic preparation); PREP
(Preparation)

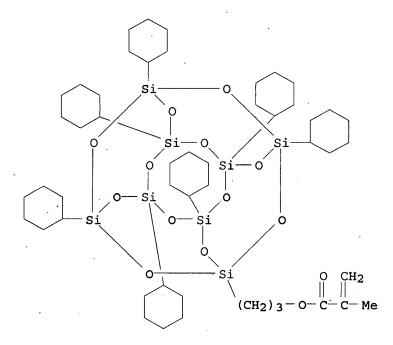
(homopolymer; preparation of methacrylate-functionalized polyhedral oligomeric silsesquioxane monomers and polymers)

RN 169699-56-3 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-(heptacyclohexylpentacyclo[9.5.1.13,9.15,15 .17,13]octasiloxanyl)propyl ester, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 169391-90-6 CMF C49 H88 O14 Si8



RN 169699-57-4 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,1 5.17,13]octasiloxanyl)propyl ester, homopolymer (9CI) (CA INDEX NAME)

CM 1

IT 169391-90-6P 169391-91-7P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP

(Preparation); RACT (Reactant or reagent)

(macromer; preparation of methacrylate-functionalized polyhedral oligomeric silsesquioxane monomers and polymers)

RN 169391-90-6 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-(heptacyclohexylpentacyclo[9.5.1.13,9.15,15.17,13]octasiloxanyl)propyl ester (9CI) (CA INDEX NAME)

RN 169391-91-7 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-(heptacyclopentylpentacyclo[9.5.1.13,9.15,1

KATHLEEN FULLER EIC1700 571/272-2505

5.17,13]octasiloxanyl)propyl ester (9CI) (CA INDEX NAME)